

# DESCRIPTION

## Substituted Arylalkanoic Acid Derivative and Use Thereof

### Field of the Invention

The present invention relates to a novel substituted arylalkanoic acid derivative. More specifically, the present invention relates to a substituted arylalkanoic acid derivative having an action as a medicament and a synthetic intermediate of said compound.

### Background Art

Various kinds of prostaglandins and various kinds of leukotrienes are produced in the bodies of mammals in response to variety of stimuli such as inflammatory stimuli and physical stimuli.

Both of prostaglandins and leukotrienes are metabolites of arachidonic acid, and they are physiologically active substances referred to as lipid mediators, and they cause various physiological responses of mammals by binding to receptors expressed on surfaces of various cells or in the cells.

Arachidonic acid is produced from a phospholipid as a substrate, such as phosphatidylcholine which is a cell membrane component, with the aid of an enzymatic activity of phospholipase A<sub>2</sub> (PLA<sub>2</sub>).

Arachidonic acid produced by the action of PLA<sub>2</sub> is converted into prostaglandin (PG) H<sub>2</sub> with the aid of an enzymatic activity of constitutive type cyclooxygenase (COX) 1 or inducible type COX-2, and further converted into PGE<sub>2</sub>, PGD<sub>2</sub>, PGF<sub>2</sub>α, PGI<sub>2</sub>, thromboxane (TX) A<sub>2</sub> and the like with the aid of each synthetic enzyme. Further, arachidonic acid is also metabolized by the action of 5-lipoxygenase (5-LO) and thereby converted to leukotriene (LT) A<sub>4</sub>, and further

converted to LTB<sub>4</sub>, LTC<sub>4</sub>, LTD<sub>4</sub>, LTE<sub>4</sub> and the like by the enzymatic activities of LTA<sub>4</sub> hydrolase, LTC<sub>4</sub> synthase, glutathione S-transferase and the like [Goodman & Gilman, Pharmacological Basis of Therapeutics, 9th edition, p.801, 1999 (Hirokawa Shoten); Funk, C.D., SCIENCE, vol. 294, p.1871, 2001].

Prostaglandins bind to each specific receptor to cause inflammatory reactions such as fervescence, enhancement of vascular permeability, vasodilation, swelling and pain, bronchial smooth muscle contraction, platelet aggregation, tumor cell proliferation, enhancement of bone resorption, nerve cell degeneration and the like, and thus play important roles in expression of symptoms or pathological formation for various diseases.

Leukotrienes are physiologically active substances which bind to each specific receptor to cause inflammatory reactions such as excessive accumulation of leucocytes and enhancement of vascular permeability, smooth muscle contraction, mucus secretion, proliferation of tumor cells and the like, and thus play important roles in expression of symptoms or pathological formation for various diseases.

Although inflammatory reactions themselves are essential reactions for living bodies to survive when they face pathogenic substances and affections, they are sometimes excessively caused or continue without any reason for providing evident benefit under certain situations or in certain diseases [Goodman & Gilman, Pharmacological Basis of Therapeutics, 9th edition, p.827, 1999 (Hirokawa Shoten)]. The condition of living body referred to in this specification wherein an acute or chronic inflammatory reaction is observed means a condition that an excessive or unbeneficial acute or temporary inflammatory reaction or chronic and persistent inflammatory reaction is caused. Further, an inflammatory reaction refers to a series of events caused by stimuli, for example, physical hazards such as heat, infective substances, ischemia, antigen/antibody reaction and the like, and it is accompanied by flare, swelling, hyperalgesia, algescic onset and the like as well-



known macroscopic clinical symptoms. It is known that, as histological mechanisms for these reactions, vasodilation, enhancement of vascular permeability, infiltration of leucocytes and phagocytes, histological decomposition and fibrosing and the like are caused [Goodman & Gilman, Pharmacological Basis of Therapeutics, 9th edition, p.827, 1999 (Hirokawa Shoten)]. It is known that many of these histological reactions are caused by prostaglandins and/or leukotrienes, and prostaglandins and/or leukotrienes plays important roles in inflammatory reactions.

For example, it was reported that, in a pathological tissue of rheumatoid arthritis, which is an autoimmune and chronic inflammatory disease, expression of COX-2 and production of PGE<sub>2</sub> and TXA<sub>2</sub> as well as expression of 5-LO and production of LTB<sub>4</sub> were observed (Bonnet et al., Prostaglandins, 1995, vol. 50, p.127), and in a mouse deficient in FLAP, which is a protein required for activation of 5-LO, symptom of collagen-induced arthritis, which is a pathological model of chronic rheumatoid arthritis, was milder compared with that in a wild-type mouse (Griffiths et al., J. Exp. Med., 1997, vol. 185, p.1123), and thus it has been suggested that prostaglandins and leukotrienes play important roles in the pathological formation of chronic rheumatoid arthritis.

It was reported that, in a pathological tissue of bronchial asthma, one of chronic allergic diseases, excessive production of PGD<sub>2</sub> and TXA<sub>2</sub> as well as excessive production of LTC<sub>4</sub> and LTD<sub>4</sub> were observed (Wenzel et al., Am Rev. Respir. Dis., 1990, vol. 142, p.112), and an airway hypersensitive reaction, which is a pathological model of bronchial asthma, was unlikely to occur in a PGD<sub>2</sub> receptor-deficient mouse (Matsuoka et al., SCIENCE, vol. 287, p.2013, 2000). Thus, it has been demonstrated that roles of prostaglandins and leukotrienes are important in bronchial asthma.

In a cerebral tissue after ischemic reperfusion, expression of COX-2

increased, and concentrations of PGE<sub>2</sub> and TXA<sub>2</sub> increased, whereas activity of 5-LO increased, and production amount of LTC<sub>4</sub> increased (Ohtsuki et al., Am. J. Physiol., 1995, vol. 268, p.1249). Thus, it is known that prostaglandins and leukotrienes play important roles in formation of infarct that is accepted as an ischemic reperfusion injury.

It has been revealed that, in a pathological tissue of Alzheimer's disease, one of the diseases with neurodegeneration, the COX activity and 5-LO activity increased, prostaglandins and leukotrienes cause formation of the  $\beta$ -amyloid protein, one of the pathogenic substances of Alzheimer's disease, and further cause degeneration of nerve cells (Sugaya et al., Jpn. J. Pharmacol., 2000, vol. 82, p.85), and thus it is believed that prostaglandins and leukotrienes play important roles in the formation of neurodegenerative diseases such as Alzheimer's disease.

Furthermore, for example, it was reported that, in a pathological tissue of colon cancer, COX and 5-LO were expressed, and amounts of production of prostaglandins and leukotrienes were increased (Dreyling et al., Biochim. Biophys. Acta., 1986, vol. 878, p.184), and leukotriene caused increase in colon cancer cells (Qiao et al., Biochim. Biophys. Acta, 1995, vol. 1258, p.215; Hong et al., Cancer Res., 1999, vol. 59, p.2223). Thus, it is believed that prostaglandins and leukotrienes play important roles also in tissues of large bowel cancer.

Involvement of prostaglandins and/or leukotrienes in diseases and pathological conditions is not limited to those diseases exemplified above, and it has been demonstrated that prostaglandins and/or leukotrienes are involved in variety of conditions, various diseases, or various pathological conditions where acute or chronic inflammatory reactions are observed and their roles are important.

For the above reason, various prostaglandin production suppressors or leukotriene production suppressors are used as agents for prophylactic or therapeutic treatment of conditions, various diseases or pathological conditions

where an acute or chronic inflammatory reaction is recognized. Various non-steroidal anti-inflammatory drugs (NSAIDS) as medicaments having a prostaglandin production-suppressing action are available and used as therapeutic agents for chronic rheumatoid arthritis and osteoarthritis, antiphlogistic analgesic agents for injury and the like, prophylactic agents for cerebral infarction or myocardial infarction, prophylactic agents for colon polyposis and the like. However, the class of NSAIDS suppress only production of prostaglandins, and as a result, they increase amounts of production of leukotrienes, and exhibit side effects such as asthmatic attack and gastrointestinal injury as well as renal disturbance. Furthermore, a difference between an effective dose and a dose inducing the side effects is small in these NSAIDS, and no satisfactory agent is available from a viewpoint of therapeutic effect. A 5-LO inhibitor is available which is described in European Patent No. 279263 as a medicament having a leukotriene production-suppressing action, and the inhibitor is known as a prophylactic agent for asthma. However, since the agent causes side effects such as hepatic disorder, its dosage is limited, and the agent is not satisfactory also from a viewpoint of therapeutic effect. Since steroid agents suppress production of both of prostaglandins and leukotrienes, they are used as prophylactic agents or therapeutic agents for conditions of living bodies, various diseases and pathological conditions where various acute or chronic inflammatory reactions are observed. However, their actions are not limited to the lipid mediator production-suppressing action, and they exhibit severe side effects such as induction and exacerbation of infectious diseases due to the immunosuppression action, growth retardation due to normal cell antiproliferative activity, anetoderma and peptic ulcer. Therefore, their uses are limited.

Furthermore, for the above reasons, it is considered that compounds, that suppress the production of both of prostaglandins and leukotrienes and have reduced side effect, are effective as therapeutic agents or prophylactic agents for

such conditions of living bodies, diseases or pathological conditions in mammals as described above, and methods of using such compounds together with medicaments available at present are more effective therapeutic or prophylactic methods.

Therefore, development of compounds suppressing the production of both of prostaglandins and leukotrienes, and manufacture of pharmaceutical preparations thereof are strongly desired.

As compounds structurally similar to the compounds of the present invention, for example, biphenyl-5-alkanoic acid derivatives and use thereof are described in WO99/19291. However, the moiety of these compounds that corresponds to "AR" included in the formula (I) of the compounds of the present invention is phenyl group, and thus structural features of the above compounds are different. Further, biaryl phospholipase A<sub>2</sub> inhibitors are described in U.S. Patent No. 5,391,817 [Japanese Patent Unexamined Publication (Kokai) No. 7-22399]. However, the moiety of these compounds that corresponds to "AR" included in the formula (I) of the compounds of the present invention is only defined to be phenyl group, and thus the structural features of the above compounds are different. Bicyclic heterocyclic compounds are described in WO00/35886 as protease inhibitors. However, the substituents of these compounds on the moiety that corresponds to "AR" included in the formula (I) of the compounds of the present invention are different, and further, the publication is completely silent about whether or not the compounds described in the above patent document have any prostaglandin production-suppressing action or leukotriene production-suppressing action.

[Patent document 1] WO99/19291

[Patent document 2] U.S. Patent No. 5,391,817

[Patent document 3] WO00/35886

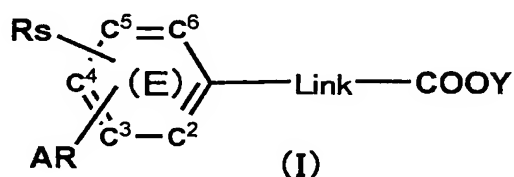
Disclosure of the Invention

An object of the present invention is to provide a novel compound having superior prostaglandin production-suppressing action and leukotriene production-suppressing action. Another object of the present invention is to provide a compound for prophylactic and/or therapeutic treatment of various inflammatory diseases, autoimmune diseases, allergic diseases, pain and fibrosis in mammals caused by lipid mediators. A further object of the present invention is to provide a pharmaceutical composition containing such a compound. A still further object of the present invention is to provide an intermediate for the production of the compound. These objects and other objects as well as advantages of the present invention will be apparent for those skilled in the art from the following descriptions.

In order to achieve the aforementioned objects, the inventors of the present invention conducted various researches. As a result, they found that the substituted arylalkanoic acid derivatives represented by the following general formula, which are novel compounds, had superior prostaglandin production-suppressing action and leukotriene production-suppressing action, and thus accomplished the present invention.

The present invention is embodied by, for example, those described in the following (1) to (191).

(1) A compound represented by the formula (I):



[In the formula, Link represents a saturated or unsaturated straight hydrocarbon chain having 1 to 3 carbon atoms.

C<sup>2</sup>, C<sup>3</sup>, C<sup>4</sup>, C<sup>5</sup>, and C<sup>6</sup> in the aromatic ring (E) independently represent a

ring-constituting carbon atom. One of the ring-constituting carbon atoms to which Rs and AR do not bind may be replaced with V.

V represents nitrogen atom, or carbon atom substituted with Zx. Zx represents a linear or branched saturated alkyl group having 1 to 4 carbon atoms, fluorine atom, chlorine atom, bromine atom, nitro group,  $\text{-OR}^9$ , or  $\text{-N(Rn}^1\text{)(Rn}^2\text{)}$ .  $\text{R}^9$  represents hydrogen atom, a lower alkyl group having 1 to 4 carbon atoms, or  $\text{-A}^6\text{-Qp}$ , wherein  $\text{A}^6$  represents a single bond or methylene, Qp represents phenyl group, and the phenyl group may be substituted with one of  $\text{T}^1$  or two or more of the same or different  $\text{T}^1$ .  $\text{T}^1$  represents a linear or branched saturated alkyl group having 1 to 4 carbon atoms, hydroxyl group, fluorine atom, chlorine atom, bromine atom, trifluoromethyl group, nitro group, an alkoxy group having 1 to 4 carbon atoms, or a mono- or dialkylamino group having 1 to 4 carbon atoms.  $\text{Rn}^1$  represents hydrogen atom or a linear or branched saturated alkyl group having 1 to 4 carbon atoms,  $\text{Rn}^2$  has the same meaning as  $\text{Rn}^1$ , or represents  $\text{-COR}^{23}$  or  $\text{-SO}_2\text{R}^{24}$ , or binds to  $\text{Rn}^1$  to form a 3- to 6-membered ring together with the nitrogen atom to which they bind to form a saturated nitrogen-containing cycloalkyl group or morpholino group.  $\text{R}^{23}$  represents hydrogen atom, a lower alkyl group having 1 to 4 carbon atoms, a lower alkoxy group having 1 to 4 carbon atoms,  $\text{-O-A}^6\text{-Qp}$ , or  $\text{-N(R}^{25}\text{)(R}^{26}\text{)}$ .  $\text{R}^{25}$  represents hydrogen atom, or a linear or branched saturated alkyl group having 1 to 4 carbon atoms.  $\text{R}^{26}$  has the same meaning as  $\text{R}^{25}$ , or binds to  $\text{R}^{25}$  to form a 3- to 6-membered ring together with the nitrogen atom to which they bind to form a saturated nitrogen-containing cycloalkyl group or morpholino group.  $\text{R}^{24}$  represents a lower alkyl group having 1 to 4 carbon atoms, amino group, or a mono- or dialkylamino group having 1 to 4 carbon atoms.

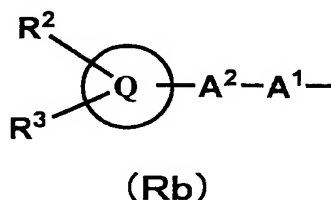
Rs represents  $\text{-D-Rx}$  or  $\text{-N(Ry)(Rz)}$ .

D represents a single bond, oxygen atom, sulfur atom,  $\text{-S(O)-}$ ,  $\text{-S(O)}_2\text{-}$ , or  $\text{-C(O)-}$ .

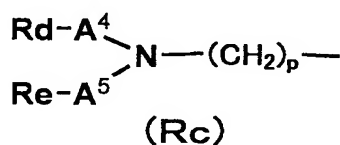
Rx represents a linear or branched saturated alkyl group having 3 to 8 carbon atoms, or represents Ra represented by the following formula:



Rb represented by the following formula:



or Rc represented by the following formula:



k in Ra represents 0 or an integer of 1 to 3. R<sup>1</sup> represents a saturated cyclic alkyl group having 3 to 7 carbon atoms, or a condensed saturated cyclic alkyl group having 6 to 8 carbon atoms, and R<sup>1</sup> may be substituted with one of lower alkyl group having 1 to 4 carbon atoms or two or more of the same or different lower alkyl groups having 1 to 4 carbon atoms. Q in Rb represents a partially unsaturated or completely unsaturated monocyclic or condensed bicyclic carbon ring or a heterocyclic ring (q), and binds to A<sup>2</sup> at an arbitrary position. The heterocyclic ring (q) contains the same or different 1 to 4 ring-constituting heteroatoms selected from the group consisting of nitrogen atom, oxygen atom, and sulfur atom. A<sup>1</sup> represents a single bond or an alkylene (a) having 1 to 3 carbon atoms, and the alkylene (a) may be substituted with a lower alkyl group having 1 to 4 carbon atoms or phenyl group. A<sup>2</sup> represents a single bond, oxygen atom, sulfur atom, -S(O)-, -S(O)<sub>2</sub>-, or -N(R<sup>4</sup>)- (provided that when A<sup>2</sup> represents oxygen atom, sulfur atom, -S(O)-, -S(O)<sub>2</sub>- or -N(R<sup>4</sup>)-, A<sup>1</sup> represents ethylene or trimethylene). R<sup>2</sup> and R<sup>3</sup> independently represent hydrogen atom, a linear or branched saturated

alkyl group having 1 to 4 carbon atoms, oxo group, thioxo group, fluorine atom, chlorine atom, bromine atom, trifluoromethyl group,  $-OR^5$ ,  $-N(R^6)(R^6')$ ,  $-NHCOR^7$ ,  $-NHSO_2R^8$ , or  $-A^6-Qa$ , or they bind to each other to represent methylenedioxy group.  $Qa$  represents a partially unsaturated or completely unsaturated monocyclic or condensed bicyclic carbon ring or a heterocyclic ring ( $qa$ ), binds to  $A^6$  at an arbitrary position on the ring, and may be substituted with one of  $T^1$  or two or more of the same or different  $T^1$ . The heterocyclic ring ( $qa$ ) contains the same or different 1 to 4 ring-constituting heteroatoms selected from the group consisting of nitrogen atom, oxygen atom, and sulfur atom.  $R^4$  and  $R^6$  independently represent hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms.  $R^5$  and  $R^7$  independently represent hydrogen atom, a lower alkyl group having 1 to 4 carbon atoms, or  $-A^6-Qa$ .  $R^8$  represents a lower alkyl group having 1 to 4 carbon atoms.  $R^6'$  has the same meaning as  $R^6$ , or binds to  $R^6$  to form a 3- to 6-membered ring together with the nitrogen atom to which they bind to represent a saturated nitrogen-containing cycloalkyl group or morpholino group.  $p$  in  $R_c$  represents an integer of 2 to 4.  $A^4$  represents a single bond, methylene, or ethylene.  $A^5$  represents  $-C(O)-$ ,  $-C(S)-$ , or  $-S(O)_2-$ .  $R_d$  represents hydrogen atom, an alkyl group having 1 to 8 carbon atoms, or  $Qa$ .  $R_e$  represents an alkyl group having 1 to 8 carbon atoms,  $-A^6-Qa$ ,  $-(CH_2)_iR^{14}$ ,  $-OR^{28}$ ,  $-SR^{28}$ , or  $-N(R^{29})(R^{30})$ .  $i$  represents an integer of 1 to 3,  $R^{14}$  represents hydroxyl group, an alkoxy group having 1 to 4 carbon atoms, carboxyl group, or an N,N-dialkylcarbamoyl group having 1 to 4 carbon atoms.  $R^{28}$  represents an alkyl group having 1 to 8 carbon atoms, or  $-A^6-Qa$ .  $R^{29}$  represents an alkyl group having 1 to 8 carbon atoms, an alkoxycarbonyl group having 1 to 4 carbon atoms, or  $-A^6-Qa$ .  $R^{30}$  represents hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms, or binds to  $R^{29}$  to form a 3- to 6-membered ring together with the nitrogen atom to which they bind to represent a saturated nitrogen-containing cycloalkyl group or morpholino group.



Rz has the same meaning as Rx, or Rz represents methyl group, ethyl group, or -A<sup>5</sup>-Re. Ry represents hydrogen atom, an alkyl group having 1 to 8 carbon atoms, or -A<sup>6</sup>-Qp, or Ry may bind to Rz to form, together with a nitrogen atom to which they bind, a saturated or unsaturated 3 to 7-membered nitrogen-containing cyclic group, wherein said nitrogen-containing cyclic group may optionally be substituted with one or two lower alkyl groups having 1 to 4 carbon atoms wherein said two alkyl groups may be the same or different.

AR represents a partially unsaturated or completely unsaturated condensed bicyclic carbon ring or a heterocyclic ring (ar), and may be substituted with one of Xa or two or more of the same or different Xa. The heterocyclic ring (ar) contains the same or different 1 to 4 ring-constituting heteroatoms selected from the group consisting of nitrogen atom, oxygen atom, and sulfur atom. Xa represents a linear or branched saturated alkyl group having 1 to 4 carbon atoms, a saturated cyclic alkyl group having 3 to 7 carbon atoms, oxo group, thioxo group, fluorine atom, chlorine atom, trifluoromethyl group, -(CH<sub>2</sub>)<sub>i</sub>R<sup>14</sup>, -OR<sup>10</sup>, -N(R<sup>11</sup>)(R<sup>12</sup>), -SO<sub>2</sub>R<sup>13</sup>, or -COR<sup>27</sup>. R<sup>10</sup> represents hydrogen atom, a lower alkyl group having 1 to 4 carbon atoms, or -(CH<sub>2</sub>)<sub>i</sub>R<sup>14</sup>. R<sup>11</sup> represents hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms. R<sup>12</sup> represents hydrogen atom, a lower alkyl group having 1 to 4 carbon atoms, a hydroxyalkyl group having 2 to 4 carbon atoms, -COR<sup>15</sup>, or -SO<sub>2</sub>R<sup>16</sup>, or binds to R<sup>11</sup> to form a 3- to 6-membered ring together with the nitrogen atom to which they bind to represent a saturated nitrogen-containing cycloalkyl group or morpholino group. R<sup>15</sup> represents a lower alkyl group having 1 to 4 carbon atoms, a hydroxyalkyl group having 2 to 4 carbon atoms, amino group, a mono- or dialkylamino group having 1 to 4 carbon atoms, or -A<sup>6</sup>-Qa. R<sup>13</sup> and R<sup>16</sup> independently represent a lower alkyl group having 1 to 4 carbon atoms, amino group, or a mono- or dialkylamino group having 1 to 4 carbon atoms. R<sup>27</sup> represents hydrogen atom, hydroxyl group, an alkoxy group having 1 to 4 carbon

atoms, a lower alkyl group having 1 to 4 carbon atoms, amino group, or a mono- or dialkylamino group having 1 to 4 carbon atoms.

Y represents hydrogen atom, a lower alkyl group having 1 to 4 carbon atoms,  $-(CH_2)_mN(R^{18})(R^{19})$ , or  $-C(R^{20})_2OC(O)A^3R^{21}$ . Symbol m represents an integer of 2 or 3.  $R^{18}$  is the same as  $R^{19}$ , or binds to  $R^{19}$  to form a 3- to 6-membered ring together with the nitrogen atom to which they bind to represent a saturated nitrogen-containing cycloalkyl group or morpholino group.  $R^{19}$  represents methyl group, ethyl group, or propyl group.  $R^{20}$  represents hydrogen atom, methyl group, ethyl group, or propyl group.  $R^{21}$  represents a lower alkyl group having 1 to 4 carbon atoms, a cyclic saturated alkyl group having 3 to 6 carbon atoms, or phenyl group, and  $A^3$  represents a single bond, or oxygen atom. This compound may sometimes be hereinafter referred to simply as "Compound (I)" of the present invention."] or a salt thereof.

(1-2) The compound or salt thereof according to (1), wherein, in the formula (I), Link is  $-(CH_2)_n$ , n is an integer of 1 to 3,  $R_z$  has the same meaning as that of  $R_x$  or represents  $-A^5-Re$  when  $R_s$  is  $-N(R_y)(R_z)$ , and  $R_y$  is hydrogen atom, an alkyl group having 1 to 8 carbon atoms, or  $A^6-Qp$ , or  $R_y$  binds to  $R_z$  to form, together with a nitrogen atom to which they bind, a saturated or unsaturated 3 to 7-membered nitrogen-containing cyclic group.

(2) The compound or salt thereof according to (1) or (1-2) mentioned above, wherein, in the formula (I), AR binds to any atom among  $C^2$  and  $C^3$  in the aromatic ring (E).

(3) The compound or salt thereof according to any one of (1) to (2) mentioned above, wherein, in the formula (I), n is an integer of 2 (the description of "according to any one of (1) to (2)" includes (1-2) mentioned above, and the same or similar description should be construed in the same manner hereinafter in the specification).

(4) The compound or salt thereof according to any one of (1) to (3) mentioned above,

wherein, in the formula (I), AR is a residue of naphthalene, benzofuran, benzo[b]thiophene, indole, benzothiazole, dihydro-3H-benzothiazole, quinoline, dihydro-1H-quinoline, benzo[d]isothiazole, 1H-indazole, benzo[c]isothiazole, 2H-indazole, imidazo[1,2-a]pyridine, 1H-pyrrolo[2,3-b]pyridine, isoquinoline, dihydro-2H-isoquinoline, cinnoline, quinazoline, quinoxaline, 1H-benzimidazole, benzoxazole, 1H-pyrrolo[3,2-b]pyridine, benzo[1,2,5]thiadiazole, 1H-benzotriazole, 1,3-dihydropyrrolo[2,3-b]pyridine, 1,3-dihydrobenzimidazole, dihydro-3H-benzoxazole, phthalazine, [1,8]naphthalidine, [1,5]naphthalidine, 1H-pyrrolo[3,2-c]pyridine, 1H-pyrrolo[2,3-c]pyridine, 1H-pyrazolo[4,3-b]pyridine, 1H-pyrazolo[4,3-c]pyridine, 1H-pyrazolo[3,4-c]pyridine, 1H-pyrazolo[3,4-b]pyridine, [1,2,4]triazolo[4,3-a]pyridine, thieno[3,2-c]pyridine, thieno[3,2-b]pyridine, 1H-thieno[3,2-c]pyrazole, benzo[d]isoxazole, benzo[c]isoxazole, indolizine, 1,3-dihydroindole, 1H-pyrazolo[3,4-d]thiazole, 2H-isoindole, [1,2,4]triazolo[1,5-a]pyrimidine, 1H-pyrazolo[3,4-b]pyrazine, 1H-imidazo[4,5-b]pyrazine, 7H-purine, or 4H-chromene (the aforementioned residue may be substituted with one of Xa or two or more of the same or different Xa).

(5) The compound or salt thereof according to any one of (1) to (3) mentioned above, wherein, in the formula (I), AR is naphthalen-2-yl group, naphthalen-1-yl group, benzofuran-5-yl group, benzofuran-4-yl group, benzofuran-2-yl group, benzo[b]thiophen-5-yl group, benzo[b]thiophen-4-yl group, benzo[b]thiophen-2-yl group, indol-5-yl group, indol-4-yl group, indol-6-yl group, benzothiazol-6-yl group, benzothiazol-7-yl group, benzothiazol-5-yl group, benzothiazol-4-yl group, dihydro-3H-benzothiazol-6-yl group, dihydro-3H-benzothiazol-7-yl group, dihydro-3H-benzothiazol-5-yl group, dihydro-3H-benzothiazol-4-yl group, quinolin-6-yl group, quinolin-3-yl group, quinolin-5-yl group, quinolin-7-yl group, dihydro-1H-quinolin-6-yl group, dihydro-1H-quinolin-5-yl group, benzo[d]isothiazol-5-yl group, benzo[d]isothiazol-4-yl group, benzo[d]isothiazol-6-yl group, benzo[d]isothiazol-7-yl

group, 1H-indazol-5-yl group, 1H-indazol-4-yl group, 1H-indazol-6-yl group,  
 benzo[c]isothiazol-5-yl group, benzo[c]isothiazol-4-yl group, benzo[c]isothiazol-6-yl  
 group, benzo[c]isothiazol-7-yl group, 2H-indazol-5-yl group, 2H-indazol-4-yl group,  
 2H-indazol-6-yl group, imidazo[1,2-a]pyridin-6-yl group, imidazo[1,2-a]pyridin-7-yl  
 group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1H-pyrrolo[2,3-b]pyridin-4-yl group,  
 isoquinolin-6-yl group, isoquinolin-3-yl group, isoquinolin-5-yl group, isoquinolin-7-  
 yl group, dihydro-2H-isoquinolin-6-yl group, dihydro-2H-isoquinolin-5-yl group,  
 cinnolin-6-yl group, cinnolin-5-yl group, quinazolin-6-yl group, quinazolin-7-yl  
 group, quinazolin-5-yl group, quinoxalin-2-yl group, quinoxalin-6-yl group,  
 quinoxalin-5-yl group, 1H-benzimidazol-5-yl group, 1H-benzimidazol-4-yl group,  
 benzoxazol-5-yl group, benzoxazol-6-yl group, benzoxazol-4-yl group, benzoxazol-7-  
 yl group, 1H-pyrrolo[3,2-b]pyridin-5-yl group, 1H-pyrrolo[3,2-b]pyridin-6-yl group,  
 benzo[1,2,5]thiadiazol-5-yl group, benzo[1,2,5]thiadiazol-4-yl group, 1H-  
 benzotriazol-5-yl group, 1H-benzotriazol-4-yl group, 1,3-dihydropyrrolo[2,3-  
 b]pyridin-5-yl group, 1,3-dihydropyrrolo[2,3-b]pyridin-4-yl group, 1,3-  
 dihydrobenzimidazol-5-yl group, 1,3-dihydrobenzimidazol-4-yl group, dihydro-3H-  
 benzoxazol-6-yl group, dihydro-3H-benzoxazol-7-yl group, dihydro-3H-benzoxazol-5-  
 yl group, dihydro-3H-benzoxazol-4-yl group, phthalazin-6-yl group, phthalazin-5-yl  
 group, [1,8]naphthalidin-3-yl group, [1,8]naphthalidin-4-yl group,  
 [1,5]naphthalidin-3-yl group, [1,5]naphthalidin-4-yl group, 1H-pyrrolo[3,2-  
 c]pyridin-6-yl group, 1H-pyrrolo[3,2-c]pyridin-4-yl group, 1H-pyrrolo[2,3-c]pyridin-  
 5-yl group, 1H-pyrrolo[2,3-c]pyridin-4-yl group, 1H-pyrazolo[4,3-b]pyridin-5-yl  
 group, 1H-pyrazolo[4,3-b]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-6-yl group,  
 1H-pyrazolo[4,3-c]pyridin-4-yl group, 1H-pyrazolo[3,4-c]pyridin-5-yl group, 1H-  
 pyrazolo[3,4-c]pyridin-4-yl group, 1H-pyrazolo[3,4-b]pyridin-5-yl group, 1H-  
 pyrazolo[3,4-b]pyridin-4-yl group, [1,2,4]triazolo[4,3-a]pyridin-6-yl group,  
 [1,2,4]triazolo[4,3-a]pyridin-7-yl group, thieno[3,2-c]pyridin-2-yl group, thieno[3,2-

c]pyridin-3-yl group, thieno[3,2-c]pyridin-6-yl group, thieno[3,2-b]pyridin-2-yl group, thieno[3,2-b]pyridin-3-yl group, thieno[3,2-b]pyridin-5-yl group, thieno[3,2-b]pyridin-6-yl group, 1H-thieno[3,2-c]pyrazol-5-yl group, 1H-thieno[3,2-c]pyrazol-4-yl group, benzo[d]isoxazol-5-yl group, benzo[d]isoxazol-4-yl group, benzo[d]isoxazol-6-yl group, benzo[d]isoxazol-7-yl group, benzo[c]isoxazol-5-yl group, benzo[c]isoxazol-4-yl group, benzo[c]isoxazol-6-yl group, benzo[c]isoxazol-7-yl group, indolizin-7-yl group, indolizin-6-yl group, indolizine-8-yl group, 1,3-dihydroindol-5-yl group, 1,3-dihydroindol-4-yl group, 1,3-dihydroindol-6-yl group, 1H-pyrazolo[3,4-d]thiazol-5-yl group, 2H-isoindol-5-yl group, 2H-isoindol-4-yl group, [1,2,4]triazolo[1,5-a]pyrimidin-6-yl group, 1H-pyrazolo[3,4-b]pyrazin-5-yl group, 1H-imidazo[4,5-b]pyrazin-5-yl group, 7H-purin-2-yl group, 4H-chromen-6-yl group, or 4H-chromen-5-yl group (the aforementioned groups may be substituted with one of Xa or two or more of the same or different Xa).

(6) The compound or salt thereof according to any one of (1) to (3) mentioned above, wherein, in the formula (I), AR is a residue of naphthalene, benzofuran, benzo[b]thiophene, indole, benzothiazole, dihydro-3H-benzothiazole, quinoline, dihydro-1H-quinoline, benzo[d]isothiazole, 1H-indazole, benzo[c]isothiazole, 2H-indazole, imidazo[1,2-a]pyridine, 1H-pyrrolo[2,3-b]pyridine, isoquinoline, or dihydro-2H-isoquinoline (the aforementioned residue may be substituted with one of Xa or two or more of the same or different Xa).

(7) The compound or salt thereof according to any one of (1) to (3) mentioned above, wherein, in the formula (I), AR is a residue of cinnoline, quinazoline, quinoxaline, 1H-benzimidazole, benzoxazole, 1H-pyrrolo[3,2-b]pyridine, benzo[1,2,5]thiadiazole, 1H-benzotriazole, 1,3-dihydropyrrolo[2,3-b]pyridine, 1,3-dihydrobenzimidazole, dihydro-3H-benzoxazole, phthalazine, [1,8]naphthalidine, [1,5]naphthalidine, 1H-pyrrolo[3,2-c]pyridine, 1H-pyrrolo[2,3-c]pyridine, 1H-pyrazolo[4,3-b]pyridine, 1H-pyrazolo[4,3-c]pyridine, 1H-pyrazolo[3,4-c]pyridine, 1H-pyrazolo[3,4-b]pyridine,

[1,2,4]triazolo[4,3-a]pyridine, thieno[3,2-c]pyridine, thieno[3,2-b]pyridine, 1H-thieno[3,2-c]pyrazole, benzo[d]isoxazole, benzo[c]isoxazole, indolizine, 1,3-dihydroindole, 1H-pyrazolo[3,4-d]thiazole, 1H-pyrazolo[3,4-d]thiazole, 2H-isoindole, [1,2,4]triazolo[1,5-a]pyrimidine, 1H-pyrazolo[3,4-b]pyrazine, 1H-imidazo[4,5-b]pyrazine, 7H-purine, or 4H-chromene (the aforementioned residue may have one of Xa or two or more of the same or different Xa).

(8) The compound or salt thereof according to any one of (1) to (7) mentioned above, wherein, in the formula (I), Rs is -D-Rx or -N(Ry)(Rz), D is a single bond, oxygen atom, sulfur atom, -S(O)-, -S(O)<sub>2</sub>-, or -C(O)-, Rx is a linear or branched saturated alkyl group having 3 to 8 carbon atoms, or Ra, Rb, or Rc, k in Ra is 0 or an integer of 1 to 3, R<sup>1</sup> is a saturated cyclic alkyl group having 3 to 7 carbon atoms or a condensed saturated cyclic alkyl group having 6 to 8 carbon atoms, R<sup>1</sup> may be substituted with one of lower alkyl group having 1 to 4 carbon atoms or two or more of the same or different lower alkyl groups having 1 to 4 carbon atoms, Q in Rb is phenyl group, thienyl group, furyl group, pyrrolyl group, pyridyl group, oxazolyl group, isoxazolyl group, thiazolyl group, isothiazolyl group, imidazolyl group, pyrazolyl group, oxadiazolyl group, thiadiazolyl group, triazolyl group, tetrazolyl group, naphthyl group, tetrahydronaphthyl group, indanyl group, indenyl group, quinolyl group, isoquinolyl group, indolyl group, benzofuryl group, benzothienyl group, benzimidazolyl group, benzoxazolyl group, benzothiazolyl group, indazolyl group, 4H-chromenyl group, dihydrobenzodioxyl group, benzoisoxazolyl group, pyrrolopyridinyl group, pyrazolopyridinyl group, triazolopyridinyl group, thienopyridinyl group, thienopyrazolyl group, 1,3-dihydrobenzimidazole group, dihydro-3H-benzoxazole group, or dihydro-3H-benzothiazole group (the aforementioned groups binds to A<sup>2</sup> at an arbitrary position), A<sup>1</sup> is a single bond or an alkylene (a) having 1 to 3 carbon atoms, the alkylene (a) may be substituted with a lower alkyl group having 1 to 4 carbon atoms or phenyl group, A<sup>2</sup> is a single

bond, oxygen atom, sulfur atom,  $-S(O)-$ ,  $-S(O)_2-$ , or  $-N(R^4)-$  (provided that when  $A^2$  represents oxygen atom, sulfur atom,  $-S(O)-$ ,  $-S(O)_2-$ , or  $-N(R^4)-$ ,  $A^1$  represents ethylene or trimethylene),  $R^2$  and  $R^3$  independently represent hydrogen atom, a linear or branched saturated alkyl group having 1 to 4 carbon atoms, oxo group, thioxo group, fluorine atom, chlorine atom, bromine atom, trifluoromethyl group,  $-OR^5$ ,  $-N(R^6)(R^{6'})$ ,  $-NHCOR^7$ ,  $-NH SO_2 R^8$ , or  $-A^6-Qa$ , or they bind to each other to represent methylenedioxy group,  $Qa$  is phenyl group, pyridyl group, oxazolyl group, isoxazolyl group, thiazolyl group, isothiazolyl group, imidazolyl group, pyrazolyl group, oxadiazolyl group, thiadiazolyl group, triazolyl group, tetrazolyl group, naphthyl group, indanyl group, indenyl group, quinolyl group, isoquinolyl group, indolyl group, benzofuryl group, benzothienyl group, benzimidazolyl group, benzoxazolyl group, benzothiazolyl group, or indazolyl group (these groups may be substituted with one of  $T^1$  or two or more of the same or different  $T^1$ , and bind to  $A^6$  at an arbitrary position on the ring),  $R^4$  and  $R^6$  independently represent hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms,  $R^5$  and  $R^7$  independently represent hydrogen atom, a lower alkyl group having 1 to 4 carbon atoms, or  $-A^6-Qa$ ,  $R^8$  is a lower alkyl group having 1 to 4 carbon atoms,  $R^{6'}$  has the same meaning as  $R^6$ , or binds to  $R^6$  to form a 3- to 6-membered ring together with the nitrogen atom to which they bind to form a saturated nitrogen-containing cycloalkyl group or morpholino group,  $p$  in  $Rc$  is an integer of 2 to 4,  $A^4$  is a single bond or methylene or ethylene,  $A^5$  is  $-C(O)-$ ,  $-C(S)-$ , or  $-S(O)_2-$ ,  $Rd$  is hydrogen atom, an alkyl group having 1 to 8 carbon atoms, or  $Qa$ ,  $Re$  is an alkyl group having 1 to 8 carbon atoms,  $-A^6-Qa$ ,  $-(CH_2)_i R^{14}$ ,  $-OR^{28}$ ,  $-SR^{28}$ , or  $-N(R^{29})(R^{30})$ ,  $i$  is an integer of 1 to 3,  $R^{14}$  is hydroxyl group, an alkoxy group having 1 to 4 carbon atoms, carboxyl group, or an  $N,N$ -dialkylcarbamoyl group having 1 to 4 carbon atoms,  $R^{28}$  is an alkyl group having 1 to 8 carbon atoms or  $-A^6-Qa$ ,  $R^{29}$  is an alkyl group having 1 to 8 carbon atoms, an alkoxycarbonyl group having 1 to 4 carbon atoms, or  $-A^6-Qa$  group,  $R^{30}$  is hydrogen

atom or a lower alkyl group having 1 to 4 carbon atoms, or binds to R<sup>29</sup> to form a 3- to 6-membered ring together with the nitrogen atom to which they bind to form a saturated nitrogen-containing cycloalkyl group or morpholino group, R<sub>z</sub> has the same meaning as R<sub>x</sub>, or is -A<sup>5</sup>-R<sub>e</sub>, and R<sub>y</sub> is hydrogen atom, an alkyl group having 1 to 8 carbon atoms, or -A<sup>6</sup>-Q<sub>p</sub>, or binds to R<sub>z</sub> to form a saturated or unsaturated nitrogen-containing cyclic substituent having 3 to 7 atoms together with nitrogen atom to which they binds.

(9) The compound or salt thereof according to any one of (1) to (8) mentioned above, wherein, in the formula (I), among C<sup>2</sup>, C<sup>3</sup>, C<sup>4</sup>, C<sup>5</sup>, and C<sup>6</sup> in the aromatic ring (E), one ring-constituting atom to which R<sub>s</sub> or AR does not bind is replaced with nitrogen atom.

(10) The compound or salt thereof according to any one of (1) to (8) mentioned above, wherein, in the formula (I), among C<sup>2</sup>, C<sup>3</sup>, C<sup>4</sup>, C<sup>5</sup>, or C<sup>6</sup> in the aromatic ring (E), one ring-constituting atom to which R<sub>s</sub> or AR does not bind is replaced with -N(R<sub>n</sub><sup>1</sup>)(R<sub>n</sub><sup>2</sup>) (provided that one of R<sub>n</sub><sup>1</sup> and R<sub>n</sub><sup>2</sup> represents a substituent other than hydrogen atom).

(11) The compound or salt thereof according to (1) or (10) mentioned above, wherein, in the formula (I), R<sub>s</sub> is -O-R<sub>x</sub>.

(12) The compound or salt thereof according to any one of (1) to (11) mentioned above, wherein, in the formula (I), R<sub>s</sub> is -O-R<sub>c</sub>.

(13) The compound or salt thereof according to any one of (1) to (10) mentioned above, wherein, in the formula (I), R<sub>s</sub> is -N(R<sub>y</sub>)(R<sub>z</sub>).

(14) The compound or salt thereof according to any one of (1) to (10) mentioned above, wherein, in the formula (I), R<sub>s</sub> is -D-R<sub>x</sub>, and D is a single bond, sulfur atom, -S(O)-, -S(O)<sub>2</sub>-, or -C(O)-.

(15) The compound or salt thereof according to any one of (1) to (10) mentioned above, wherein, in the formula (I), R<sub>s</sub> is -S-R<sub>x</sub>.



(16) The compound or salt thereof according to (1-2) mentioned above, wherein, in the formula (I), AR binds at the position of C<sup>2</sup> in the aromatic ring (E), and Rs binds to one of the ring-constituting carbon atoms C<sup>3</sup>, C<sup>4</sup>, and C<sup>5</sup>.

(17) The compound or salt thereof according to (16) mentioned above, wherein, in the formula (I), Rs is -O-Rx, and no ring-constituting carbon atom in the aromatic ring (E) is replaced with V.

(18) The compound or salt thereof according to (16) or (17) mentioned above, wherein, in the formula (I), n is an integer of 2, and Y is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms.

(19) The compound or salt thereof according to (4) mentioned above, wherein, in the formula (I), Link is -(CH<sub>2</sub>)<sub>n</sub>-, n is an integer of 2, AR binds at the position of C<sup>2</sup> in the aromatic ring (E), Rs binds to one of ring-constituting carbon atoms C<sup>3</sup>, C<sup>4</sup> and C<sup>5</sup>, Rs is -O-Rx, Y is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms, and all of C<sup>2</sup>, C<sup>3</sup>, C<sup>4</sup>, C<sup>5</sup>, and C<sup>6</sup> in the aromatic ring (E) are not replaced with V.

(20) The compound or salt thereof according to (5) mentioned above, wherein, in the formula (I), Link is -(CH<sub>2</sub>)<sub>n</sub>-, n is an integer of 2, AR binds at the position of C<sup>2</sup> in the aromatic ring (E), Rs binds to one of ring-constituting carbon atoms C<sup>3</sup>, C<sup>4</sup> and C<sup>5</sup>, Rs is -O-Rx, Y is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms, and all of C<sup>2</sup>, C<sup>3</sup>, C<sup>4</sup>, C<sup>5</sup>, and C<sup>6</sup> in the aromatic ring (E) are not replaced with V.

(21) The compound or salt thereof according to any one of (16) to (20) mentioned above, wherein, in the formula (I), Xa which may substitute on AR is methyl group, ethyl group, propyl group, hydroxyethyl group, carboxymethyl group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, amino group, methylamino group, dimethylamino group, carboxyl group, carbamoyl group, acetyl group, methanesulfonyl group, sulfamoyl group, or N,N-dimethylsulfamoyl group.

(22) The compound or salt thereof according to any one of (16) to (21) mentioned above, wherein, in the formula (I),  $R_s$  is  $-O-R_x$ ,  $R_x$  is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, or cyclohexylmethyl group, or  $R_b$  (provided that  $Q$  in  $R_b$  is phenyl group or indan-2-yl group),  $A^1$  is a single bond, a methylene group substituted with methyl group or ethyl group, or unsubstituted methylene group, or an ethylene group substituted with methyl group or ethyl group, or unsubstituted ethylene group,  $A^2$  is a single bond, oxygen atom, sulfur atom,  $-N(\text{methyl})-$ , or  $-N(\text{ethyl})-$  (provided that when  $A^2$  represents oxygen atom, sulfur atom,  $-N(\text{methyl})-$ , or  $-N(\text{ethyl})-$ ,  $A^1$  represents ethylene), and  $R^2$  and  $R^3$  independently represent hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, or dimethylamino group (provided that when  $Q$  is phenyl group,  $A^1$  is a single bond or unsubstituted methylene, and  $A^2$  is a single bond, one of  $R^2$  and  $R^3$  is a substituent other than hydrogen atom).

(23) The compound or salt thereof according to any one of (16) to (22) mentioned above, wherein, in the formula (I),  $R_x-D$  binds at the position of  $C^3$  in the aromatic ring (E).

(24) The compound or salt thereof according to any one of (16) to (22) mentioned above, wherein, in the formula (I),  $R_x-D$  binds at the position of  $C^4$  in the aromatic ring (E).

(25) The compound or salt thereof according to any one of (16) to (22) mentioned above, wherein, in the formula (I),  $R_x-D$  binds at the position of  $C^5$  in the aromatic ring (E).

(26) The compound or salt thereof according to (1-2) mentioned above, wherein, in the formula (I),  $n$  is an integer of 1 to 3,  $AR$  binds to  $C^2$ ,  $R_s$  binds to one of the ring-constituting carbon atoms  $C^3$ ,  $C^4$ , and  $C^5$ , a ring-constituting atom among  $C^3$ ,  $C^4$ , and  $C^5$  to which  $R_s$  does not bind may be replaced with  $V$ ,

V is nitrogen atom or carbon atom substituted with Zx, Zx is fluorine atom, chlorine atom, bromine atom, nitro group, methyl group, hydroxyl group, methoxy group, amino group, N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, or N,N-dimethylsulfamoylamino group,

Rs is -D-Rx or -N(Ry)(Rz), D is oxygen atom or sulfur atom, Rx is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-cyclopentylethyl group, or 2-cyclohexylethyl group, or Rb or Rc, Q in Rb is phenyl group, thienyl group, furyl group, pyridyl group, oxazolyl group, naphthyl group, tetrahydronaphthyl group, indanyl group, indolyl group, or dihydrobenzodioxyl group, and A<sup>2</sup> is a single bond, oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)- (provided that when A<sup>2</sup> is oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)-, A<sup>1</sup> represents ethylene). R<sup>2</sup> and R<sup>3</sup> independently represent hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, dimethylamino group, acetylamino group, or methylsulfonylamino group (provided that when Q is phenyl group, A<sup>1</sup> is a single bond or unsubstituted methylene, and A<sup>2</sup> is a single bond, one of R<sup>2</sup> and R<sup>3</sup> is a substituent other than hydrogen atom). Symbol p in Rc is an integer of 2 or 3, A<sup>4</sup> is a single bond or methylene, A<sup>5</sup> is -C(O)-, -C(S)-, or -S(O)<sub>2</sub>-, Rd is hydrogen atom, or methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group, cyclopropylmethyl group, cyclopentyl group, cyclopentylmethyl group, cyclohexyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, or pyridin-4-yl group, Re is methyl group, ethyl group, propyl group, isopropyl group, butyl group,

isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, phenylmethyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, furan-2-yl group, furan-3-yl group, thiophen-2-yl group, thiophen-3-yl group, methoxy group, ethoxy group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenoxy group, 4-methylphenoxy group, 4-chlorophenoxy group, 4-fluorophenoxy group, thiomethoxy group, amino group, N-methylamino group, N,N-dimethylamino group, N-ethylamino group, N,N-diethylamino group, N-propylamino group, N-isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group, N-(4-fluorophenyl)amino group, N-(pyridin-2-yl)amino group, N-(pyridin-3-yl)amino group, N-(pyridin-4-yl)amino group, N-(furan-2-yl)amino group, N-(furan-3-yl)amino group, N-(thiophen-2-yl)amino group, N-(thiophen-3-yl)amino group, pyrrolidino group, piperidino group, morpholino group, methyloxycarbonylamino group, or ethyloxycarbonylamino group,

Rz is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl

group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl

group, 2-(N-phenyl-N-methylamino)ethyl group, 2-(N-ethyl-N-phenylamino)ethyl  
 group, isobutyryl group, isopropylthiocarbonyl group, isopropylsulfonyl group,  
 valeryl group, butylthiocarbonyl group, isovaleryl group, isobutylthiocarbonyl group,  
 pivaloyl group, t-butylthiocarbonyl group, cyclopropylcarbonyl group,  
 cyclopropylthiocarbonyl group, cyclopentylcarbonyl group, cyclopentylthiocarbonyl  
 group, cyclohexylcarbonyl group, cyclohexylthiocarbonyl group,  
 cyclopentylmethylcarbonyl group, cyclopentylmethylthiocarbonyl group,  
 cyclohexylmethylcarbonyl group, cyclohexylmethylthiocarbonyl group, benzoyl  
 group, thiobenzoyl group, phenylsulfonyl group, 4-methylphenylcarbonyl group, 4-  
 methylphenylthiocarbonyl group, 4-methylphenylsulfonyl group, 4-  
 chlorophenylcarbonyl group, 4-chlorophenylthiocarbonyl group, 4-  
 fluorophenylcarbonyl group, 4-fluorophenylthiocarbonyl group,  
 isopropoxy carbonyl group, N-isopropylcarbamoyl group, N-isopropylthiocarbamoyl  
 group, butyloxy carbonyl group, N-butylcarbamoyl group, N-butylthiocarbamoyl  
 group, isobutyloxy carbonyl group, N-isobutylcarbamoyl group, N-  
 isobutylthiocarbamoyl group, t-butyloxy carbonyl group, N-t-butylcarbamoyl group,  
 N-t-butylthiocarbamoyl group, cyclopropyloxy carbonyl group, N-  
 cyclopropylcarbamoyl group, N-cyclopropylthiocarbamoyl group,  
 cyclopentyloxy carbonyl group, N-cyclopentylcarbamoyl group, N-  
 cyclopentylthiocarbamoyl group, cyclohexyloxy carbonyl group, N-  
 cyclohexylcarbamoyl group, N-cyclohexylthiocarbamoyl group,  
 cyclopentylmethyloxy carbonyl group, cyclohexylmethyloxy carbonyl group,  
 phenyloxy carbonyl group, N-phenylcarbamoyl group, N-phenylthiocarbamoyl group,  
 4-methylphenyloxy carbonyl group, N-(4-methylphenyl)carbamoyl group, N-(4-  
 methylphenyl)thiocarbamoyl group, 4-chlorophenyloxy carbonyl group, N-(4-  
 chlorophenyl)carbamoyl group, N-(4-chlorophenyl)thiocarbamoyl group, 4-  
 fluorophenyloxy carbonyl group, N-(4-fluorophenyl)carbamoyl group, N-(4-

fluorophenyl)thiocarbamoyl group, (pyrrolidino-1-yl)carbonyl group, (piperidino-1-yl)carbonyl group, or (morpholino-4-yl)carbonyl group, R<sub>y</sub> is hydrogen atom, methyl group, ethyl group, or isobutyl group, or binds to R<sub>z</sub> to form pyrrolidino group, piperidino group, piperazino group, morpholino group, pyrrol-1-yl group, imidazol-1-yl group, or pyrazol-1-yl group together with nitrogen atom to which they binds,

AR is naphthalen-2-yl group, naphthalen-1-yl group, benzofuran-5-yl group, benzofuran-4-yl group, benzofuran-2-yl group, benzo[b]thiophen-5-yl group, benzo[b]thiophen-4-yl group, benzo[b]thiophen-2-yl group, indol-5-yl group, indol-4-yl group, indol-6-yl group, benzothiazol-6-yl group, benzothiazol-7-yl group, benzothiazol-5-yl group, benzothiazol-4-yl group, dihydro-3H-benzothiazol-6-yl group, dihydro-3H-benzothiazol-7-yl group, dihydro-3H-benzothiazol-5-yl group, dihydro-3H-benzothiazol-4-yl group, quinolin-6-yl group, quinolin-3-yl group, quinolin-5-yl group, quinolin-7-yl group, dihydro-1H-quinolin-6-yl group, dihydro-1H-quinolin-5-yl group, benzo[d]isothiazol-5-yl group, benzo[d]isothiazol-4-yl group, benzo[d]isothiazol-6-yl group, benzo[d]isothiazol-7-yl group, 1H-indazol-5-yl group, 1H-indazol-4-yl group, 1H-indazol-6-yl group, benzo[c]isothiazol-5-yl group, benzo[c]isothiazol-4-yl group, benzo[c]isothiazol-6-yl group, benzo[c]isothiazol-7-yl group, 2H-indazol-5-yl group, 2H-indazol-4-yl group, 2H-indazol-6-yl group, imidazo[1,2-a]pyridin-6-yl group, imidazo[1,2-a]pyridin-7-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1H-pyrrolo[2,3-b]pyridin-4-yl group, isoquinolin-6-yl group, isoquinolin-3-yl group, isoquinolin-5-yl group, isoquinolin-7-yl group, dihydro-2H-isoquinolin-6-yl group, dihydro-2H-isoquinolin-5-yl group, cinnolin-6-yl group, cinnolin-5-yl group, quinazolin-6-yl group, quinazolin-7-yl group, quinazolin-5-yl group, quinoxalin-2-yl group, quinoxalin-6-yl group, quinoxalin-5-yl group, 1H-benzimidazol-5-yl group, 1H-benzimidazol-4-yl group, benzoxazol-5-yl group, benzoxazol-6-yl group, benzoxazol-4-yl group, benzoxazol-7-yl group, 1H-pyrrolo[3,2-b]pyridin-5-yl group, 1H-pyrrolo[3,2-b]pyridin-6-yl group,

benzo[1,2,5]thiadiazol-5-yl group, benzo[1,2,5]thiadiazol-4-yl group, 1H-benzotriazol-5-yl group, 1H-benzotriazol-4-yl group, 1,3-dihydropyrrolo[2,3-b]pyridin-5-yl group, 1,3-dihydropyrrolo[2,3-b]pyridin-4-yl group, 1,3-dihydrobenzimidazol-5-yl group, 1,3-dihydrobenzimidazol-4-yl group, dihydro-3H-benzoxazol-6-yl group, dihydro-3H-benzoxazol-7-yl group, dihydro-3H-benzoxazol-5-yl group, dihydro-3H-benzoxazol-4-yl group, phthalazin-6-yl group, phthalazin-5-yl group, [1,8]naphthalidin-3-yl group, [1,8]naphthalidin-4-yl group, [1,5]naphthalidin-3-yl group, [1,5]naphthalidin-4-yl group, 1H-pyrrolo[3,2-c]pyridin-6-yl group, 1H-pyrrolo[3,2-c]pyridin-4-yl group, 1H-pyrrolo[2,3-c]pyridin-5-yl group, 1H-pyrrolo[2,3-c]pyridin-4-yl group, 1H-pyrazolo[4,3-b]pyridin-5-yl group, 1H-pyrazolo[4,3-b]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-4-yl group, 1H-pyrazolo[3,4-c]pyridin-5-yl group, 1H-pyrazolo[3,4-c]pyridin-4-yl group, 1H-pyrazolo[3,4-b]pyridin-5-yl group, 1H-pyrazolo[3,4-b]pyridin-4-yl group, [1,2,4]triazolo[4,3-a]pyridin-6-yl group, [1,2,4]triazolo[4,3-a]pyridin-7-yl group, thieno[3,2-c]pyridin-2-yl group, thieno[3,2-c]pyridin-3-yl group, thieno[3,2-c]pyridin-6-yl group, thieno[3,2-b]pyridin-2-yl group, thieno[3,2-b]pyridin-3-yl group, thieno[3,2-b]pyridin-5-yl group, thieno[3,2-b]pyridin-6-yl group, 1H-thieno[3,2-c]pyrazol-5-yl group, 1H-thieno[3,2-c]pyrazol-4-yl group, benzo[d]isoxazol-5-yl group, benzo[d]isoxazol-4-yl group, benzo[d]isoxazol-6-yl group, benzo[d]isoxazol-7-yl group, benzo[c]isoxazol-5-yl group, benzo[c]isoxazol-4-yl group, benzo[c]isoxazol-6-yl group, benzo[c]isoxazol-7-yl group, indolizin-7-yl group, indolizin-6-yl group, indolizine-8-yl group, 1,3-dihydroindol-5-yl group, 1,3-dihydroindol-4-yl group, 1,3-dihydroindol-6-yl group, 1H-pyrazolo[3,4-d]thiazol-5-yl group, 2H-isoindol-5-yl group, 2H-isoindol-4-yl group, [1,2,4]triazolo[1,5-a]pyrimidin-6-yl group, 1H-pyrazolo[3,4-b]pyrazin-5-yl group, 1H-imidazo[4,5-b]pyrazin-5-yl group, 7H-purin-2-yl group, 4H-chromen-6-yl group, or 4H-chromen-5-yl group (the aforementioned groups may be substituted with one



of Xa or two or more of the same or different Xa), Xa is oxo group, thioxo group, fluorine atom, chlorine atom, trifluoromethyl group, methyl group, ethyl group, propyl group, 2-hydroxyethyl group, carboxymethyl group, 2-carboxyethyl group, N,N-dimethylcarbamoylmethyl group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, carboxymethyloxy group, 2-carboxyethyloxy group, N,N-dimethylcarbamoylmethyloxy group, amino group, methylamino group, dimethylamino group, 2-hydroxyethylamino group, carbamoylamino group, acetylamino group, furan-2-carboxyamino group, 2-hydroxyacetylamino group, 2-aminoacetylamino group, methylsulfonylamino group, (N,N-dimethylsulfamoyl)amino group, methanesulfonyl group, sulfamoyl group, N-methylsulfamoyl group, N,N-dimethylsulfamoyl group, carboxyl group, acetyl group, carbamoyl group, or N,N-dimethylcarbamoyl group, and Y is hydrogen atom, methyl group or ethyl group.

(27) The compound or salt thereof according to (1-2) mentioned above, wherein, in the formula (I), n is an integer of 2,

C<sup>2</sup> is carbon atom to which AR binds, C<sup>3</sup> is carbon atom to which Rs binds, C<sup>4</sup> may be replaced with V, C<sup>5</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms,

V is nitrogen atom, or carbon atom substituted with Zx, Zx is fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, or N,N-dimethylamino group,

Rs is -O-Rx, Rx is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group,

5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group,  
 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group,  
 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group,  
 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl  
 group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl  
 group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-  
 chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl  
 group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-  
 methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-  
 dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl  
 group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-  
 chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl  
 group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-  
 difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-  
 dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-  
 dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-  
 dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-  
 (trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-  
 (trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-  
 methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-  
 methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-  
 methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl  
 group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-  
 chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-  
 (trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-  
 (trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group,  
 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-

chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, or 2-(N-ethyl-N-phenylamino)ethyl group,

AR is naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group, 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-

5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, or benzoxazol-5-yl group, and

Y is hydrogen atom, methyl group, or ethyl group.

(28) The compound or salt thereof according to (1-2) mentioned above, wherein, in the formula (I), n is an integer of 2,

C<sup>2</sup> is carbon atom to which AR binds, C<sup>4</sup> is carbon atom to which Rs binds, C<sup>5</sup> may be replaced with V, C<sup>3</sup> and C<sup>6</sup> represents an unsubstituted ring-constituting carbon atom,

V is nitrogen atom, or carbon atom substituted with Zx, Zx is fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, or N,N-dimethylamino group,

Rs is -O-Rx, Rx is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl

group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, or 2-(N-ethyl-N-phenylamino)ethyl group,

AR is naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-

aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group, 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-

pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, or benzoxazol-5-yl group, and

Y is hydrogen atom, methyl group, or ethyl group.

(29) The compound or salt thereof according to (1-2) mentioned above, wherein, in the formula (I), AR binds to C<sup>3</sup> in the aromatic ring (E), and Rs binds to C<sup>5</sup> or C<sup>6</sup> in the aromatic ring (E).

(30) The compound or salt thereof according to (29) mentioned above, wherein, in the formula (I), Rs is -O-Rx, and all of C<sup>2</sup>, C<sup>3</sup>, C<sup>4</sup>, C<sup>5</sup>, and C<sup>6</sup> in the aromatic ring (E) are not replaced with V.

(31) The compound or salt thereof according to (29) or (30) mentioned above, wherein, in the formula (I), n is an integer of 2, and Y is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms.

(32) The compound or salt thereof according to (4) mentioned above, wherein, in the formula (I), Link is -(CH<sub>2</sub>)<sub>n</sub>-, n is an integer of 2, AR binds to C<sup>3</sup> in the aromatic ring (E), Rs binds to the ring-constituting carbon atom C<sup>5</sup> or C<sup>6</sup> in the aromatic ring (E), Rs is -O-Rx, Y is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms, and all of C<sup>2</sup>, C<sup>3</sup>, C<sup>4</sup>, C<sup>5</sup>, and C<sup>6</sup> in the aromatic ring (E) are not replaced with V.

(33) The compound or salt thereof according to (5) mentioned above, wherein, in the formula (I), n is an integer of 2, AR binds to C<sup>3</sup> in the aromatic ring (E), Rs binds to the ring-constituting carbon atom C<sup>5</sup> or C<sup>6</sup> in the aromatic ring (E), Rs is -O-Rx, Y is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms, and all of C<sup>2</sup>, C<sup>3</sup>, C<sup>4</sup>, C<sup>5</sup>, and C<sup>6</sup> in the aromatic ring (E) are not replaced with V.

(34) The compound or salt thereof according to any one of (29) to (33) mentioned above, wherein, in the formula (I), Xa which may substitute on AR is methyl group, ethyl group, propyl group, hydroxyethyl group, carboxymethyl group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, amino group, methylamino group,

dimethylamino group, carboxyl group, carbamoyl group, acetyl group, methanesulfonyl group, sulfamoyl group, or N,N-dimethylsulfamoyl group.

(35) The compound or salt thereof according to any one of (29) to (34) mentioned above, wherein, in the formula (I), Rs is -O-Rx, Rx is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, or cyclohexylmethyl group, or Rb (provided that Q in Rb is phenyl group or indan-2-yl group), A<sup>1</sup> is a single bond, or methylene group substituted with methyl group or ethyl group or unsubstituted methylene group, or ethylene group substituted with methyl group or ethyl group or unsubstituted ethylene group, A<sup>2</sup> is a single bond, oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)- (provided that when A<sup>2</sup> is oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)-, A<sup>1</sup> is ethylene), and R<sup>2</sup> and R<sup>3</sup> independently represent hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, or dimethylamino group (provided that when Q is phenyl group, A<sup>1</sup> is a single bond or unsubstituted methylene, and A<sup>2</sup> is a single bond, one of R<sup>2</sup> and R<sup>3</sup> is a substituent other than hydrogen atom).

(36) The compound or salt thereof according to any one of (29) to (35) mentioned above, wherein, in the formula (I), Rs binds at the position of C<sup>5</sup> in the aromatic ring (E).

(37) The compound or salt thereof according to any one of (29) to (35) mentioned above, wherein, in the formula (I), Rs binds at the position of C<sup>6</sup> in the aromatic ring (E).

(38) The compound or salt thereof according to (1-2) mentioned above, wherein, in the formula (I), n is an integer of 2,

C<sup>3</sup> is carbon atom to which AR binds, C<sup>5</sup> is carbon atom to which Rs binds, C<sup>2</sup>, C<sup>4</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms,

Rs is -O-Rx, Rx is butyl group, isobutyl group, 2-ethylbutyl group,



cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl

group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, or 2-(N-ethyl-N-phenylamino)ethyl group,

AR is naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group, 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-

dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, or benzoxazol-5-yl group, and

Y is hydrogen atom, methyl group, or ethyl group.

(39) The compound or salt thereof according to (1-2) mentioned above, wherein, in the formula (I), AR binds to C<sup>3</sup> in the aromatic ring (E), Rs binds to C<sup>4</sup> in the aromatic ring (E), and C<sup>6</sup> is replaced with V.

(40) The compound or salt thereof according to (39) mentioned above, wherein, in the formula (I), n is an integer of 2, V is carbon atom substituted with Zx, Rs is -O-Rx, and Y is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms.

(41) The compound or salt thereof according to (4) mentioned above, wherein, in the formula (I), Link is -(CH<sub>2</sub>)<sub>n</sub>-, n is an integer of 2, AR binds to C<sup>3</sup> in the aromatic ring (E), Rs binds to C<sup>4</sup> in the aromatic ring (E), C<sup>6</sup> is carbon atom substituted with Zx, Rs is -O-Rx, and Y is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms.

(42) The compound or salt thereof according to (5) mentioned above, wherein, in the formula (I), Link is -(CH<sub>2</sub>)<sub>n</sub>-, n is an integer of 2, AR binds to C<sup>3</sup> in the aromatic ring (E), Rs binds to C<sup>4</sup> in the aromatic ring (E), C<sup>6</sup> is carbon atom substituted with

Zx, Rs is -O-Rx, and Y is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms.

(43) The compound or salt thereof according to any one of (39) to (42) mentioned above, wherein, in the formula (I), Xa which may substitute on AR is methyl group, ethyl group, propyl group, hydroxyethyl group, carboxymethyl group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, amino group, methylamino group, dimethylamino group, carboxyl group, carbamoyl group, acetyl group, methanesulfonyl group, sulfamoyl group, or N,N-dimethylsulfamoyl group.

(44) The compound or salt thereof according to any one of (39) to (43) mentioned above, wherein, in the formula (I), Rs is -O-Rx, Rx is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, or cyclohexylmethyl group, or Rb (provided that Q in Rb is phenyl group or indan-2-yl group), A<sup>1</sup> is a single bond, or methylene group substituted with methyl group or ethyl group or unsubstituted methylene group, or ethylene group substituted with methyl group or ethyl group or unsubstituted ethylene group, A<sup>2</sup> is a single bond, oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)- (provided that when A<sup>2</sup> is oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)-, A<sup>1</sup> is ethylene), R<sup>2</sup> and R<sup>3</sup> independently represent hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, or dimethylamino group (provided that when Q is phenyl group, A<sup>1</sup> is a single bond or unsubstituted methylene, and A<sup>2</sup> is a single bond, one of R<sup>2</sup> and R<sup>3</sup> is a substituent other than hydrogen atom).

(45) The compound or salt thereof according to (1-2) mentioned above, wherein, in the formula (I), n is an integer of 2,

C<sup>3</sup> is carbon atom to which AR binds, C<sup>4</sup> is carbon atom to which Rs binds, C<sup>6</sup> is carbon atom substituted with Zx, C<sup>2</sup> and C<sup>5</sup> are unsubstituted ring-constituting carbon atoms,

Zx is fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, or N,N-dimethylamino group,

Rs is -O-Rx, Rx is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-

methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, or 2-(N-ethyl-N-phenylamino)ethyl group,

AR is naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-

hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group, 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, or benzoxazol-5-yl group, and

Y is hydrogen atom, methyl group, or ethyl group.

(46) The compound or salt thereof according to (1-2) mentioned above, wherein, in the formula (I), AR binds to C<sup>3</sup> in the aromatic ring (E), Rs binds to C<sup>4</sup> in the aromatic ring (E), C<sup>5</sup> is nitrogen atom, and C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms.

(47) The compound or salt thereof according to (46) mentioned above, wherein, in the formula (I), n is an integer of 2, D is oxygen atom, and Y is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms.

(48) The compound or salt thereof according to (4) mentioned above, wherein, in the formula (I), Link is  $\cdot(\text{CH}_2)_n\cdot$ , n is an integer of 2, AR binds to C<sup>3</sup> in the aromatic ring (E), Rs binds to C<sup>4</sup> in the aromatic ring (E), C<sup>5</sup> is nitrogen atom, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms, Rs is  $\cdot\text{O}\cdot\text{Rx}$ , and Y is hydrogen atom

or a lower alkyl group having 1 to 4 carbon atoms.

(49) The compound or salt thereof according to (5) mentioned above, wherein, in the formula (I), Link is  $-(CH_2)_n-$ ,  $n$  is an integer of 2, AR binds to  $C^3$  in the aromatic ring (E),  $R_s$  binds to  $C^4$  in the aromatic ring (E),  $C^5$  is nitrogen atom,  $C^2$  and  $C^6$  are unsubstituted ring-constituting carbon atoms,  $R_s$  is  $-O-R_x$ , and  $Y$  is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms.

(50) The compound or salt thereof according to any one of (46) to (49) mentioned above, wherein, in the formula (I),  $X_a$  which may substitute on AR is methyl group, ethyl group, propyl group, hydroxyethyl group, carboxymethyl group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, amino group, methylamino group, dimethylamino group, carboxyl group, carbamoyl group, acetyl group, methanesulfonyl group, sulfamoyl group, or  $N,N$ -dimethylsulfamoyl group.

(51) The compound or salt thereof according to any one of (46) to (50) mentioned above, wherein, in the formula (I),  $R_s$  is  $-O-R_x$ ,  $R_x$  is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, or cyclohexylmethyl group, or  $R_b$  (provided that  $Q$  in  $R_b$  is phenyl group or indan-2-yl group),  $A^1$  is a single bond, or methylene group substituted with methyl group or ethyl group or unsubstituted methylene group, or ethylene group substituted with methyl group or ethyl group or unsubstituted ethylene group,  $A^2$  is a single bond, oxygen atom, sulfur atom,  $-N(\text{methyl})-$  or  $-N(\text{ethyl})-$  (provided that when  $A^2$  is oxygen atom, sulfur atom,  $-N(\text{methyl})-$ , or  $-N(\text{ethyl})-$ ,  $A^1$  is ethylene), and  $R^2$  and  $R^3$  independently represent hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, or dimethylamino group (provided that when  $Q$  is phenyl group,  $A^1$  is a single bond or unsubstituted methylene, and  $A^2$  is a single bond, one of  $R^2$  and  $R^3$  is a substituent other than hydrogen atom).

(52) The compound or salt thereof according to (1-2) mentioned above, wherein, in



the formula (I), n is an integer of 2,

C<sup>3</sup> is carbon atom to which AR binds, C<sup>4</sup> is carbon atom to which Rs binds, C<sup>5</sup> is nitrogen atom, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms,

Rs is -O-Rx, Rx is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-

(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, or 2-(N-ethyl-N-phenylamino)ethyl group,

AR is naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-

indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group, 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, or benzoxazol-5-yl group, and

Y is hydrogen atom, methyl group, or ethyl group.

(53) The compound or salt thereof according to (1-2) mentioned above, wherein, in the formula (I), AR binds to C<sup>3</sup> in the aromatic ring (E), Rs binds to C<sup>4</sup> in the aromatic ring (E), C<sup>5</sup> is a ring-constituting carbon atom substituted with Zx, or an unsubstituted ring-constituting carbon atom, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms, Rs is -D-Rx, and D is a single bond, sulfur atom, -S(O)-, -S(O)<sub>2</sub>-, or -C(O)-.

(53-2) The compound of salt thereof according to (1-2) mentioned above, wherein, in the formula (I), AR binds to C<sup>3</sup> in the aromatic ring (E), Rs binds to C<sup>4</sup> in the aromatic ring (E), C<sup>5</sup> is nitrogen atom, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms, Rs is -D-Rx, and D is a single bond, sulfur atom, -S(O)-, -

S(O)<sub>2</sub><sup>-</sup>, or -C(O)<sup>-</sup>.

(53-3) The compound or salt thereof according to (53) or (53-2) mentioned above, wherein, in the formula (I), Rs is -D-Rx and D is single bond.

(54) The compound or salt thereof according to any one of (53) to (53-3) mentioned above, wherein, in the formula (I), n is an integer of 2, and Y is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms.

(55) The compound or salt thereof according to (4) mentioned above, wherein, in the formula (I), Link is -(CH<sub>2</sub>)<sub>n</sub><sup>-</sup>, n is an integer of 2, AR binds to C<sup>3</sup> in the aromatic ring (E), Rs binds to C<sup>4</sup> in the aromatic ring (E), C<sup>5</sup> is a ring-constituting carbon atom substituted with Zx, or an unsubstituted ring-constituting carbon atom, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms, Rs is -D-Rx, D is a single bond, sulfur atom, -S(O)<sup>-</sup>, -S(O)<sub>2</sub><sup>-</sup>, or -C(O)<sup>-</sup>, and Y is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms.

(55-2) The compound or a salt thereof according to (4) mentioned above, wherein, in the formula (I), Link is -(CH<sub>2</sub>)<sub>n</sub><sup>-</sup>, n is an integer of 2, AR binds to C<sup>3</sup> in the aromatic ring (E), Rs binds to C<sup>4</sup> in the aromatic ring (E), C<sup>5</sup> may be replaced with V, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms, Rs is -D-Rx, D is a single bond, and Y is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms.

(56) The compound or salt thereof according to (5) mentioned above, wherein, in the formula (I), Link is -(CH<sub>2</sub>)<sub>n</sub><sup>-</sup>, n is an integer of 2, AR binds to C<sup>3</sup> in the aromatic ring (E), Rs binds to C<sup>4</sup> in the aromatic ring (E), C<sup>5</sup> is a ring-constituting carbon atom substituted with Zx, or an unsubstituted ring-constituting carbon atom, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms, Rs is -D-Rx, D is a single bond, sulfur atom, -S(O)<sup>-</sup>, -S(O)<sub>2</sub><sup>-</sup>, or -C(O)<sup>-</sup>, and Y is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms.

(56-2) The compound or a salt thereof according to (5) mentioned above, wherein, in the formula (I), Link is -(CH<sub>2</sub>)<sub>n</sub><sup>-</sup>, n is an integer of 2, AR binds to C<sup>3</sup> in the aromatic

ring (E), Rs binds to C<sup>4</sup> in the aromatic ring (E), C<sup>5</sup> may be replaced with V, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms, Rs is -D-Rx, D is a single bond, and Y is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms.

(57) The compound or salt thereof according to any one of (53) to (56-2) mentioned above, wherein, in the formula (I), Xa which may substitute on AR is methyl group, ethyl group, propyl group, hydroxyethyl group, carboxymethyl group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, amino group, methylamino group, dimethylamino group, carboxyl group, carbamoyl group, acetyl group, methanesulfonyl group, sulfamoyl group, or N,N-dimethylsulfamoyl group.

(58) The compound or salt thereof according to any one of (53) to (57) mentioned above, wherein, in the formula (I), Rs is -D-Rx, Rx is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, or cyclohexylmethyl group, or Rb (provided that Q in Rb is phenyl group or indan-2-yl group), A<sup>1</sup> is a single bond, or methylene group substituted with methyl group or ethyl group or unsubstituted methylene group, or ethylene group substituted with methyl group or ethyl group or unsubstituted ethylene group, A<sup>2</sup> is a single bond, oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)- (provided that when A<sup>2</sup> represents oxygen atom, sulfur atom, -N(methyl)- or -N(ethyl)-, A<sup>1</sup> represents ethylene), and R<sup>2</sup> and R<sup>3</sup> independently represent hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, or dimethylamino group (provided that when Q is phenyl group, A<sup>1</sup> is a single bond or unsubstituted methylene, and A<sup>2</sup> is a single bond, one of R<sup>2</sup> and R<sup>3</sup> is a substituent other than hydrogen atom).

(58-2) The compound or salt thereof according to (1-2) mentioned above, wherein, in the formula (I), n is an integer of 1 to 3, AR binds to C<sup>3</sup> in the aromatic ring (E), Rs binds to C<sup>4</sup> in the aromatic ring (E), C<sup>5</sup> may be replaced with V, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms, V is nitrogen atom or V is carbon

atom substituted with Zx, Zx is any one of fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, or N,N-dimethylamino group,

Rs is -D-Rx, D is a single bond, Rx is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-cyclopentylethyl group, or 2-cyclohexylethyl group, or Rx is Rb or Rc (provided that Q in Rb is phenyl group, thienyl group, furyl group, pyridyl group, oxazolyl group, naphthyl group, tetrahydronaphthyl group, indanyl group, indolyl group, or dihydrobenzodioxyl group), A<sup>2</sup> is a single bond, oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)- (provided that when A<sup>2</sup> represents oxygen atom, sulfur atom, -N(methyl)- or -N(ethyl)-, A<sup>1</sup> represents ethylene), R<sup>2</sup> and R<sup>3</sup> independently represent hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, dimethylamino group, acetylamino group, or methylsulfonylamino group, (provided that when Q is phenyl group, A<sup>1</sup> is a single bond or unsubstituted methylene, and A<sup>2</sup> is a single bond, one of R<sup>2</sup> and R<sup>3</sup> is a substituent other than hydrogen atom). p in Rc is an integer of 2 or 3, A<sup>4</sup> is a single bond or methylene, A<sup>5</sup> is -C(O)-, -C(S)-, or -S(O)<sub>2</sub>-, Rd is hydrogen atom, or methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group, cyclopropylmethyl group, cyclopentyl group, cyclopentylmethyl group, cyclohexyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, or pyridin-4-yl group, Re is methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, phenylmethyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group,

pyridin-3-yl group, pyridin-4-yl group, methoxy group, ethoxy group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenyloxy group, 4-methylphenyloxy group, 4-chlorophenyloxy group, 4-fluorophenyloxy group, thiomethoxy group, amino group, N-methylamino group, N,N-dimethylamino group, N-ethylamino group, N,N-diethylamino group, N-propylamino group, N-isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group, N-(4-fluorophenyl)amino group, N-(pyridin-2-yl)amino group, N-(pyridin-3-yl)amino group, N-(pyridin-4-yl)amino group, N-(furan-2-yl)amino group, N-(furan-3-yl)amino group, N-(thiophen-2-yl)amino group, N-(thiophen-3-yl)amino group, pyrrolidino group, piperidino group, morpholino group, methyloxycarbonylamino group, or ethyloxycarbonylamino group,

AR is naphthalen-2-yl group, naphthalen-1-yl group, benzofuran-5-yl group, benzofuran-4-yl group, benzofuran-2-yl group, benzo[b]thiophen-5-yl group, benzo[b]thiophen-4-yl group, benzo[b]thiophen-2-yl group, indol-5-yl group, indol-4-yl group, indol-6-yl group, benzothiazol-6-yl group, benzothiazol-7-yl group, benzothiazol-5-yl group, benzothiazol-4-yl group, dihydro-3H-benzothiazol-6-yl group, dihydro-3H-benzothiazol-7-yl group, dihydro-3H-benzothiazol-5-yl group, dihydro-3H-benzothiazol-4-yl group, quinolin-6-yl group, quinolin-3-yl group, quinolin-5-yl group, quinolin-7-yl group, dihydro-1H-quinolin-6-yl group, dihydro-1H-quinolin-5-yl group, benzo[d]isothiazol-5-yl group, benzo[d]isothiazol-4-yl group, benzo[d]isothiazol-6-yl group, benzo[d]isothiazol-7-yl group, 1H-indazol-5-yl group, 1H-indazol-4-yl group, 1H-indazol-6-yl group, benzo[c]isothiazol-5-yl group, benzo[c]isothiazol-4-yl group, benzo[c]isothiazol-6-yl group, benzo[c]isothiazol-7-yl

group, 2H-indazol-5-yl group, 2H-indazol-4-yl group, 2H-indazol-6-yl group,  
 imidazo[1,2-a]pyridin-6-yl group, imidazo[1,2-a]pyridin-7-yl group, 1H-pyrrolo[2,3-  
 b]pyridin-5-yl group, 1H-pyrrolo[2,3-b]pyridin-4-yl group, isoquinolin-6-yl group,  
 isoquinolin-3-yl group, isoquinolin-5-yl group, isoquinolin-7-yl group, dihydro-2H-  
 isoquinolin-6-yl group, dihydro-2H-isoquinolin-5-yl group, cinnolin-6-yl group,  
 cinnolin-5-yl group, quinazolin-6-yl group, quinazolin-7-yl group, quinazolin-5-yl  
 group, quinoxalin-2-yl group, quinoxalin-6-yl group, quinoxalin-5-yl group, 1H-  
 benzimidazol-5-yl group, 1H-benzimidazol-4-yl group, benzoxazol-5-yl group,  
 benzoxazol-6-yl group, benzoxazol-4-yl group, benzoxazol-7-yl group, 1H-  
 pyrrolo[3,2-b]pyridin-5-yl group, 1H-pyrrolo[3,2-b]pyridin-6-yl group,  
 benzo[1,2,5]thiadiazol-5-yl group, benzo[1,2,5]thiadiazol-4-yl group, 1H-  
 benzotriazol-5-yl group, 1H-benzotriazol-4-yl group, 1,3-dihydropyrrolo[2,3-  
 b]pyridin-5-yl group, 1,3-dihydropyrrolo[2,3-b]pyridin-4-yl group, 1,3-  
 dihydrobenzimidazol-5-yl group, 1,3-dihydrobenzimidazol-4-yl group, dihydro-3H-  
 benzoxazol-6-yl group, dihydro-3H-benzoxazol-7-yl group, dihydro-3H-benzoxazol-5-  
 yl group, dihydro-3H-benzoxazol-4-yl group, phthalazin-6-yl group, phthalazin-5-yl  
 group, [1,8]naphthalidin-3-yl group, [1,8]naphthalidin-4-yl group,  
 [1,5]naphthalidin-3-yl group, [1,5]naphthalidin-4-yl group, 1H-pyrrolo[3,2-  
 c]pyridin-6-yl group, 1H-pyrrolo[3,2-c]pyridin-4-yl group, 1H-pyrrolo[2,3-c]pyridin-  
 5-yl group, 1H-pyrrolo[2,3-c]pyridin-4-yl group, 1H-pyrazolo[4,3-b]pyridin-5-yl  
 group, 1H-pyrazolo[4,3-b]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-6-yl group,  
 1H-pyrazolo[4,3-c]pyridin-4-yl group, 1H-pyrazolo[3,4-c]pyridin-5-yl group, 1H-  
 pyrazolo[3,4-c]pyridin-4-yl group, 1H-pyrazolo[3,4-b]pyridin-5-yl group, 1H-  
 pyrazolo[3,4-b]pyridin-4-yl group, [1,2,4]triazolo[4,3-a]pyridin-6-yl group,  
 [1,2,4]triazolo[4,3-a]pyridin-7-yl group, thieno[3,2-c]pyridin-2-yl group, thieno[3,2-  
 c]pyridin-3-yl group, thieno[3,2-c]pyridin-6-yl group, thieno[3,2-b]pyridin-2-yl group,  
 thieno[3,2-b]pyridin-3-yl group, thieno[3,2-b]pyridin-5-yl group, thieno[3,2-



b]pyridin-6-yl group, 1H-thieno[3,2-c]pyrazol-5-yl group, 1H-thieno[3,2-c]pyrazol-4-yl group, benzo[d]isoxazol-5-yl group, benzo[d]isoxazol-4-yl group, benzo[d]isoxazol-6-yl group, benzo[d]isoxazol-7-yl group, benzo[c]isoxazol-5-yl group, benzo[c]isoxazol-4-yl group, benzo[c]isoxazol-6-yl group, benzo[c]isoxazol-7-yl group, indolizin-7-yl group, indolizin-6-yl group, indolizine-8-yl group, 1,3-dihydroindol-5-yl group, 1,3-dihydroindol-4-yl group, 1,3-dihydroindol-6-yl group, 1H-pyrazolo[3,4-d]thiazol-5-yl group, 2H-isoindol-5-yl group, 2H-isoindol-4-yl group, [1,2,4]triazolo[1,5-a]pyrimidin-6-yl group, 1H-pyrazolo[3,4-b]pyrazin-5-yl group, 1H-imidazo[4,5-b]pyrazin-5-yl group, 7H-purin-2-yl group, 4H-chromen-6-yl group, or 4H-chromen-5-yl group (the aforementioned groups may be substituted with one or more of Xa or two or more of the same or different Xa), Xa is oxo group, thioxo group, fluorine atom, chlorine atom, trifluoromethyl group, methyl group, ethyl group, propyl group, 2-hydroxyethyl group, carboxymethyl group, 2-carboxyethyl group, N,N-dimethylcarbamoylmethyl group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, carboxymethyloxy group, 2-carboxyethyloxy group, N,N-dimethylcarbamoylmethyloxy group, amino group, methylamino group, dimethylamino group, 2-hydroxyethylamino group, carbamoylamino group, acetylamino group, furan-2-carboxyamino group, 2-hydroxyacetylamino group, 2-aminoacetylamino group, methylsulfonylamino group, (N,N-dimethylsulfamoyl)amino group, methanesulfonyl group, sulfamoyl group, N-methylsulfamoyl group, N,N-dimethylsulfamoyl group, carboxyl group, acetyl group, carbamoyl group, or N,N-dimethylcarbamoyl group, and

Y is hydrogen atom, methyl group, or ethyl group.

(58-3) The compound or salt thereof according to (1-2) mentioned above, wherein, in the formula (I), n is an integer of 2, AR binds to C<sup>3</sup> in the aromatic ring (E), R<sub>s</sub> binds to C<sup>4</sup> in the aromatic ring (E), C<sup>5</sup> may be replaced with V, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms, V is nitrogen atom or V is carbon

atom substituted with Zx, Zx is any one of fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, or N,N-dimethylamino group,

Rs is -D-Rx, D is a single bond, Rx is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 2-methylphenyl group, 3-methylphenyl group, 4-methylphenyl group, 2,3-dimethylphenyl group, 3,5-dimethylphenyl group, 2-methoxyphenyl group, 3-methoxyphenyl group, 4-methoxyphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, 2,3-difluorophenyl group, 2,4-difluorophenyl group, 2,5-difluorophenyl group, 3,4-difluorophenyl group, 2,3-dichlorophenyl group, 2,4-dichlorophenyl group, 2,5-dichlorophenyl group, 2,6-dichlorophenyl group, 3,4-dichlorophenyl group, 3,5-dichlorophenyl group, 2-trifluoromethylphenyl group, 3-trifluoromethylphenyl group, 4-trifluoromethylphenyl group, 4-(N,N-dimethylamino)phenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, furan-2-yl group, furan-3-yl group, thiophen-2-yl group, thiophen-3-yl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, naphthalen-1-yl group, naphthalen-2-yl group, 1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, biphenyl-2-yl group, biphenyl-3-yl group, biphenyl-4-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-

methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, or 2-(N-ethyl-N-phenylamino)ethyl group,

AR is naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-

methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group,  
 benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-  
 methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-  
 indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-  
 dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-  
 5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group,  
 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-  
 methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-  
 indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-  
 indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-  
 indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-  
 hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-  
 indol-5-yl group, benzothiazol-6-yl group, 2-methylbenzothiazol-6-yl group, 2-  
 methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-  
 dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-  
 dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-  
 dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-  
 methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-  
 5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl  
 group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-  
 yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-  
 methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl  
 group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-  
 pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-  
 dihydroisoquinolin-6-yl group, cinnolin-6-yl group, or benzoxazol-5-yl group, and

Y is hydrogen atom, methyl group, or ethyl group.

(58-4) The compound or salt thereof according to (1-2) mentioned above, wherein, in the formula (I),  $n$  is an integer of 2, AR binds to C<sup>3</sup> in the aromatic ring (E), R<sub>s</sub> binds to C<sup>4</sup> in the aromatic ring (E), C<sup>5</sup> may be replaced with V, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms,

V is nitrogen atom or V is carbon atom substituted with Z<sub>x</sub>, Z<sub>x</sub> is any one of fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, or N,N-dimethylamino group,

R<sub>s</sub> is -D-R<sub>x</sub>, D is a single bond, R<sub>x</sub> is phenyl group, 2-methylphenyl group, 3-methylphenyl group, 4-methylphenyl group, 2,3-dimethylphenyl group, 3,5-dimethylphenyl group, 2-methoxyphenyl group, 3-methoxyphenyl group, 4-methoxyphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, 2,3-difluorophenyl group, 2,4-difluorophenyl group, 2,5-difluorophenyl group, 3,4-difluorophenyl group, 2,3-dichlorophenyl group, 2,4-dichlorophenyl group, 2,5-dichlorophenyl group, 2,6-dichlorophenyl group, 3,4-dichlorophenyl group, 3,5-dichlorophenyl group, 2-trifluoromethylphenyl group, 3-trifluoromethylphenyl group, 4-trifluoromethylphenyl group, 4-(N,N-dimethylamino)phenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, furan-2-yl group, furan-3-yl group, thiophen-2-yl group, thiophen-3-yl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, naphthalen-1-yl group, naphthalen-2-yl group, 1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1H-indazol-5-yl group, or 1-methyl-1H-indazol-5-yl group,

AR is naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethoxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group, 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-

methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, or benzoxazol-5-yl group, and

Y is hydrogen atom, methyl group, or ethyl group.

(58-5) The compound or salt thereof according to (1-2) mentioned above, wherein, in the formula (I), n is an integer of 2, AR binds to C<sup>3</sup> in the aromatic ring (E),

Rs binds to C<sup>4</sup> in the aromatic ring (E), C<sup>2</sup>, C<sup>5</sup>, and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms, Rs is -D-Rx, D is a single bond, Rx is phenyl group, 2-methylphenyl group, 3-methylphenyl group, 4-methylphenyl group, 2,3-dimethylphenyl group, 3,5-dimethylphenyl group, 2-methoxyphenyl group, 3-methoxyphenyl group, 4-methoxyphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, 2,3-difluorophenyl group, 2,4-difluorophenyl group, 2,5-difluorophenyl group, 3,4-difluorophenyl group, 2,3-dichlorophenyl group, 2,4-dichlorophenyl group, 2,5-dichlorophenyl group, 2,6-dichlorophenyl group, 3,4-dichlorophenyl group, 3,5-dichlorophenyl group, 2-trifluoromethylphenyl group, 3-trifluoromethylphenyl group, 4-trifluoromethylphenyl group, 4-(N,N-dimethylamino)phenyl group, indan-2-yl group, furan-2-yl group, furan-3-yl group, thiophen-2-yl group, thiophen-3-yl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, naphthalen-1-yl group, naphthalen-2-yl group, 1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1H-indazol-5-yl group, or 1-methyl-1H-indazol-5-yl group,

AR is naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethoxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl

group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group, 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, or benzoxazol-5-yl group, and Y



is hydrogen atom, methyl group, or ethyl group.

(59) The compound or salt thereof according to (1-2) mentioned above, wherein, in the formula (I), n is an integer of 1 to 3,

C<sup>3</sup> is carbon atom to which AR binds, C<sup>4</sup> is carbon atom to which Rs binds, C<sup>5</sup> may be replaced with V, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms,

V is nitrogen atom, or carbon atom substituted with Zx, Zx is fluorine atom, chlorine atom, bromine atom, nitro group, methyl group, hydroxyl group, methoxy group, amino group, N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, or N,N-dimethylsulfamoylamino group,

Rs is -S-Rx, Rx is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-cyclopentylethyl group, or 2-cyclohexylethyl group, or Rb or Rc, Q in Rb is phenyl group, thienyl group, furyl group, pyridyl group, oxazolyl group, naphthyl group, tetrahydronaphthyl group, indanyl group, indolyl group, or dihydrobenzodioxyl group, A<sup>2</sup> is a single bond, oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)- (provided that when A<sup>2</sup> is oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)-, A<sup>1</sup> is ethylene), R<sup>2</sup> and R<sup>3</sup> independently represent hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, dimethylamino group, acetylamino group, or methylsulfonylamino group (provided that when Q is phenyl group, A<sup>1</sup> is a single bond or unsubstituted methylene, and A<sup>2</sup> is a single bond, one of R<sup>2</sup> and R<sup>3</sup> is a substituent other than hydrogen atom), p in Rc is an integer of 2 or 3, A<sup>4</sup> is a single bond or methylene, A<sup>5</sup> is -C(O)-, -C(S)-, or -S(O)<sub>2</sub>-, Rd is hydrogen atom, or methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group,

cyclopropylmethyl group, cyclopentyl group, cyclopentylmethyl group, cyclohexyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, or pyridin-4-yl group, Re is methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, phenylmethyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, methoxy group, ethoxy group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenyloxy group, 4-methylphenyloxy group, 4-chlorophenyloxy group, 4-fluorophenyloxy group, thiomethoxy group, amino group, N-methylamino group, N,N-dimethylamino group, N-ethylamino group, N,N-diethylamino group, N-propylamino group, N-isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group, N-(4-fluorophenyl)amino group, N-(pyridin-2-yl)amino group, N-(pyridin-3-yl)amino group, N-(pyridin-4-yl)amino group, N-(furan-2-yl)amino group, N-(furan-3-yl)amino group, N-(thiophen-2-yl)amino group, N-(thiophen-3-yl)amino group, pyrrolidino group, piperidino group, morpholino group, methyloxycarbonylamino group, or ethyloxycarbonylamino group,

AR is naphthalen-2-yl group, naphthalen-1-yl group, benzofuran-5-yl group, benzofuran-4-yl group, benzofuran-2-yl group, benzo[b]thiophen-5-yl group, benzo[b]thiophen-4-yl group, benzo[b]thiophen-2-yl group, indol-5-yl group, indol-4-

yl group, indol-6-yl group, benzothiazol-6-yl group, benzothiazol-7-yl group,  
 benzothiazol-5-yl group, benzothiazol-4-yl group, dihydro-3H-benzothiazol-6-yl  
 group, dihydro-3H-benzothiazol-7-yl group, dihydro-3H-benzothiazol-5-yl group,  
 dihydro-3H-benzothiazol-4-yl group, quinolin-6-yl group, quinolin-3-yl group,  
 quinolin-5-yl group, quinolin-7-yl group, dihydro-1H-quinolin-6-yl group, dihydro-  
 1H-quinolin-5-yl group, benzo[d]isothiazol-5-yl group, benzo[d]isothiazol-4-yl group,  
 benzo[d]isothiazol-6-yl group, benzo[d]isothiazol-7-yl group, 1H-indazol-5-yl group,  
 1H-indazol-4-yl group, 1H-indazol-6-yl group, benzo[c]isothiazol-5-yl group,  
 benzo[c]isothiazol-4-yl group, benzo[c]isothiazol-6-yl group, benzo[c]isothiazol-7-yl  
 group, 2H-indazol-5-yl group, 2H-indazol-4-yl group, 2H-indazol-6-yl group,  
 imidazo[1,2-a]pyridin-6-yl group, imidazo[1,2-a]pyridin-7-yl group, 1H-pyrrolo[2,3-  
 b]pyridin-5-yl group, 1H-pyrrolo[2,3-b]pyridin-4-yl group, isoquinolin-6-yl group,  
 isoquinolin-3-yl group, isoquinolin-5-yl group, isoquinolin-7-yl group, dihydro-2H-  
 isoquinolin-6-yl group, dihydro-2H-isoquinolin-5-yl group, cinnolin-6-yl group,  
 cinnolin-5-yl group, quinazolin-6-yl group, quinazolin-7-yl group, quinazolin-5-yl  
 group, quinoxalin-2-yl group, quinoxalin-6-yl group, quinoxalin-5-yl group, 1H-  
 benzimidazol-5-yl group, 1H-benzimidazol-4-yl group, benzoxazol-5-yl group,  
 benzoxazol-6-yl group, benzoxazol-4-yl group, benzoxazol-7-yl group, 1H-  
 pyrrolo[3,2-b]pyridin-5-yl group, 1H-pyrrolo[3,2-b]pyridin-6-yl group,  
 benzo[1,2,5]thiadiazol-5-yl group, benzo[1,2,5]thiadiazol-4-yl group, 1H-  
 benzotriazol-5-yl group, 1H-benzotriazol-4-yl group, 1,3-dihydropyrrolo[2,3-  
 b]pyridin-5-yl group, 1,3-dihydropyrrolo[2,3-b]pyridin-4-yl group, 1,3-  
 dihydrobenzimidazol-5-yl group, 1,3-dihydrobenzimidazol-4-yl group, dihydro-3H-  
 benzoxazol-6-yl group, dihydro-3H-benzoxazol-7-yl group, dihydro-3H-benzoxazol-5-  
 yl group, dihydro-3H-benzoxazol-4-yl group, phthalazin-6-yl group, phthalazin-5-yl  
 group, [1,8]naphthalidin-3-yl group, [1,8]naphthalidin-4-yl group,  
 [1,5]naphthalidin-3-yl group, [1,5]naphthalidin-4-yl group, 1H-pyrrolo[3,2-

c]pyridin-6-yl group, 1H-pyrrolo[3,2-c]pyridin-4-yl group, 1H-pyrrolo[2,3-c]pyridin-5-yl group, 1H-pyrrolo[2,3-c]pyridin-4-yl group, 1H-pyrazolo[4,3-b]pyridin-5-yl group, 1H-pyrazolo[4,3-b]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-4-yl group, 1H-pyrazolo[3,4-c]pyridin-5-yl group, 1H-pyrazolo[3,4-c]pyridin-4-yl group, 1H-pyrazolo[3,4-b]pyridin-5-yl group, 1H-pyrazolo[3,4-b]pyridin-4-yl group, [1,2,4]triazolo[4,3-a]pyridin-6-yl group, [1,2,4]triazolo[4,3-a]pyridin-7-yl group, thieno[3,2-c]pyridin-2-yl group, thieno[3,2-c]pyridin-3-yl group, thieno[3,2-c]pyridin-6-yl group, thieno[3,2-b]pyridin-2-yl group, thieno[3,2-b]pyridin-3-yl group, thieno[3,2-b]pyridin-5-yl group, thieno[3,2-b]pyridin-6-yl group, 1H-thieno[3,2-c]pyrazol-5-yl group, 1H-thieno[3,2-c]pyrazol-4-yl group, benzo[d]isoxazol-5-yl group, benzo[d]isoxazol-4-yl group, benzo[d]isoxazol-6-yl group, benzo[d]isoxazol-7-yl group, benzo[c]isoxazol-5-yl group, benzo[c]isoxazol-4-yl group, benzo[c]isoxazol-6-yl group, benzo[c]isoxazol-7-yl group, indolizin-7-yl group, indolizin-6-yl group, indolizine-8-yl group, 1,3-dihydroindol-5-yl group, 1,3-dihydroindol-4-yl group, 1,3-dihydroindol-6-yl group, 1H-pyrazolo[3,4-d]thiazol-5-yl group, 2H-isoindol-5-yl group, 2H-isoindol-4-yl group, [1,2,4]triazolo[1,5-a]pyrimidin-6-yl group, 1H-pyrazolo[3,4-b]pyrazin-5-yl group, 1H-imidazo[4,5-b]pyrazin-5-yl group, 7H-purin-2-yl group, 4H-chromen-6-yl group, or 4H-chromen-5-yl group (the aforementioned groups may be substituted with one of Xa or two or more of the same or different Xa), Xa is oxo group, thioxo group, fluorine atom, chlorine atom, trifluoromethyl group, methyl group, ethyl group, propyl group, 2-hydroxyethyl group, carboxymethyl group, 2-carboxyethyl group, N,N-dimethylcarbamoylmethyl group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, carboxymethyloxy group, 2-carboxyethyloxy group, N,N-dimethylcarbamoylmethyloxy group, amino group, methylamino group, dimethylamino group, 2-hydroxyethylamino group, carbamoylamino group, acetylamino group, furan-2-carboxyamino group, 2-hydroxyacetylamino group, 2-

aminoacetyl amino group, methylsulfonylamino group, (N,N-dimethylsulfamoyl) amino group, methanesulfonyl group, sulfamoyl group, N-methylsulfamoyl group, N,N-dimethylsulfamoyl group, carboxyl group, acetyl group, carbamoyl group, or N,N-dimethylcarbamoyl group, and

Y is hydrogen atom, methyl group, or ethyl group.

(59-2) The compound or salt thereof according to (1) mentioned above, wherein, in the formula (I), AR binds to C<sup>3</sup> in the aromatic ring (E), Rs binds to C<sup>4</sup> in the aromatic ring (E), C<sup>5</sup> is a ring-constituting carbon atom substituted with Zx, or an unsubstituted ring-constituting carbon atom, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms, and Rs is -N(Ry)(Rz).

(59-3) The compound or salt thereof according to (1) mentioned above, wherein, in the formula (I), AR binds to C<sup>3</sup> in the aromatic ring (E), Rs binds to C<sup>4</sup> in the aromatic ring (E), C<sup>5</sup> is nitrogen atom, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms, and Rs is -N(Ry)(Rz).

(60) The compound or salt thereof according to (59-2) or (59-3) mentioned above, wherein, in the formula (I), Link is -(CH<sub>2</sub>)<sub>n</sub>-, n is an integer of 2, and Y is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms.

(61) The compound or salt thereof according to (4) mentioned above, wherein, in the formula (I), Link is -(CH<sub>2</sub>)<sub>n</sub>-, n is an integer of 2, AR binds to C<sup>3</sup> in the aromatic ring (E), Rs binds to C<sup>4</sup> in the aromatic ring (E), C<sup>5</sup> is a ring-constituting carbon atom substituted with Zx, or an unsubstituted ring-constituting carbon atom, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms, Rs is -N(Ry)(Rz), and Y is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms.

(61-2) The compound or salt thereof according to (4) mentioned above, wherein, in the formula (I), Link is -(CH<sub>2</sub>)<sub>n</sub>-, n is an integer of 2, AR binds to C<sup>3</sup> in the aromatic ring (E), Rs binds to C<sup>4</sup> in the aromatic ring (E), C<sup>5</sup> is nitrogen atom, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms, Rs is -N(Ry)(Rz), and Y is hydrogen

atom or a lower alkyl group having 1 to 4 carbon atoms.

(62) The compound or salt thereof according to (5) mentioned above, wherein, in the formula (I), Link is  $-(CH_2)_n-$ ,  $n$  is an integer of 2, AR binds to  $C^3$  in the aromatic ring (E),  $R_s$  binds to  $C^4$  in the aromatic ring (E),  $C^5$  is a ring-constituting carbon atom substituted with  $Zx$ , or an unsubstituted ring-constituting carbon atom,  $C^2$  and  $C^6$  are unsubstituted ring-constituting carbon atoms,  $R_s$  is  $-N(R_y)(R_z)$ , and  $Y$  is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms.

(62-2) The compound or salt thereof according to (5) mentioned above, wherein, in the formula (I), Link is  $-(CH_2)_n-$ ,  $n$  is an integer of 2, AR binds to  $C^3$  in the aromatic ring (E),  $R_s$  binds to  $C^4$  in the aromatic ring (E),  $C^5$  is nitrogen atom,  $C^2$  and  $C^6$  are unsubstituted ring-constituting carbon atoms,  $R_s$  is  $-N(R_y)(R_z)$ , and  $Y$  is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms.

(63) The compound or salt thereof according to any one of (59-2) to (62-2) mentioned above, wherein, in the formula (I),  $X_a$  which may substitute on AR is methyl group, ethyl group, propyl group, hydroxyethyl group, carboxymethyl group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, amino group, methylamino group, dimethylamino group, carboxyl group, carbamoyl group, acetyl group, methanesulfonyl group, sulfamoyl group, or  $N,N$ -dimethylsulfamoyl group.

(64) The compound or salt thereof according to (1-2) mentioned above, wherein, in the formula (I),  $n$  is an integer of 1 to 3,  $C^3$  is carbon atom to which AR binds,  $C^4$  is carbon atom to which  $R_s$  binds,  $C^2$ ,  $C^5$ , and  $C^6$  are unsubstituted ring-constituting carbon atoms,

$R_s$  is  $-N(R_y)(R_z)$ ,  $R_z$  is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-

methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group,  
 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group,  
 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group,  
 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group,  
 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl  
 group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl  
 group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-  
 chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl  
 group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-  
 methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-  
 dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl  
 group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-  
 chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl  
 group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-  
 difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-  
 dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-  
 dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-  
 dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-  
 (trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-  
 (trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-  
 methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-  
 methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-  
 methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl  
 group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-  
 chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-  
 (trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-  
 (trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group,

2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, 2-(N-ethyl-N-phenylamino)ethyl group, isobutyryl group, isopropylthiocarbonyl group, isopropylsulfonyl group, valeryl group, butylthiocarbonyl group, isovaleryl group, isobutylthiocarbonyl group, pivaloyl group, t-butylthiocarbonyl group, cyclopropylcarbonyl group, cyclopropylthiocarbonyl group, cyclopentylcarbonyl group, cyclopentylthiocarbonyl group, cyclohexylcarbonyl group, cyclohexylthiocarbonyl group, cyclopentylmethylcarbonyl group, cyclopentylmethylthiocarbonyl group, cyclohexylmethylcarbonyl group, cyclohexylmethylthiocarbonyl group, benzoyl group, thiobenzoyl group, phenylsulfonyl group, 4-methylphenylcarbonyl group, 4-methylphenylthiocarbonyl group, 4-methylphenylsulfonyl group, 4-chlorophenylcarbonyl group, 4-chlorophenylthiocarbonyl group, 4-fluorophenylcarbonyl group, 4-fluorophenylthiocarbonyl group, isopropoxy carbonyl group, N-isopropylcarbamoyl group, N-isopropylthiocarbamoyl group, butyloxy carbonyl group, N-butylcarbamoyl group, N-butylthiocarbamoyl group, isobutyloxy carbonyl group, N-isobutylcarbamoyl group, N-isobutylthiocarbamoyl group, t-butyloxy carbonyl group, N-t-butylcarbamoyl group, N-t-butylthiocarbamoyl group, cyclopropyloxy carbonyl group, N-cyclopropylcarbamoyl group, N-cyclopropylthiocarbamoyl group, cyclopentyloxy carbonyl group, N-cyclopentylcarbamoyl group, N-cyclopentylthiocarbamoyl group, cyclohexyloxy carbonyl group, N-cyclohexylcarbamoyl group, N-cyclohexylthiocarbamoyl group, cyclopentylmethyloxy carbonyl group, cyclohexylmethyloxy carbonyl group, phenyloxy carbonyl group, N-phenylcarbamoyl group, N-phenylthiocarbamoyl group, 4-methylphenyloxy carbonyl group, N-(4-methylphenyl)carbamoyl group, N-(4-methylphenyl)thiocarbamoyl group, 4-chlorophenyloxy carbonyl group, N-(4-



chlorophenyl)carbamoyl group, N-(4-chlorophenyl)thiocarbamoyl group, 4-fluorophenyloxycarbonyl group, N-(4-fluorophenyl)carbamoyl group, N-(4-fluorophenyl)thiocarbamoyl group, (pyrrolidino-1-yl)carbonyl group, (piperidino-1-yl)carbonyl group, or (morpholino-4-yl)carbonyl group, Ry is hydrogen atom, methyl group, ethyl group or isobutyl group, or binds to Rz to form pyrrolidino group, piperidino group, piperazino group, morpholino group, pyrrol-1-yl group, imidazol-1-yl group, or pyrazol-1-yl group together with the nitrogen atom to which they bind,

AR is naphthalen-2-yl group, naphthalen-1-yl group, benzofuran-5-yl group, benzofuran-4-yl group, benzofuran-2-yl group, benzo[b]thiophen-5-yl group, benzo[b]thiophen-4-yl group, benzo[b]thiophen-2-yl group, indol-5-yl group, indol-4-yl group, indol-6-yl group, benzothiazol-6-yl group, benzothiazol-7-yl group, benzothiazol-5-yl group, benzothiazol-4-yl group, dihydro-3H-benzothiazol-6-yl group, dihydro-3H-benzothiazol-7-yl group, dihydro-3H-benzothiazol-5-yl group, dihydro-3H-benzothiazol-4-yl group, quinolin-6-yl group, quinolin-3-yl group, quinolin-5-yl group, quinolin-7-yl group, dihydro-1H-quinolin-6-yl group, dihydro-1H-quinolin-5-yl group, benzo[d]isothiazol-5-yl group, benzo[d]isothiazol-4-yl group, benzo[d]isothiazol-6-yl group, benzo[d]isothiazol-7-yl group, 1H-indazol-5-yl group, 1H-indazol-4-yl group, 1H-indazol-6-yl group, benzo[c]isothiazol-5-yl group, benzo[c]isothiazol-4-yl group, benzo[c]isothiazol-6-yl group, benzo[c]isothiazol-7-yl group, 2H-indazol-5-yl group, 2H-indazol-4-yl group, 2H-indazol-6-yl group, imidazo[1,2-a]pyridin-6-yl group, imidazo[1,2-a]pyridin-7-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1H-pyrrolo[2,3-b]pyridin-4-yl group, isoquinolin-6-yl group, isoquinolin-3-yl group, isoquinolin-5-yl group, isoquinolin-7-yl group, dihydro-2H-isoquinolin-6-yl group, dihydro-2H-isoquinolin-5-yl group, cinnolin-6-yl group, cinnolin-5-yl group, quinazolin-6-yl group, quinazolin-7-yl group, quinazolin-5-yl group, quinoxalin-2-yl group, quinoxalin-6-yl group, quinoxalin-5-yl group, 1H-

benzimidazol-5-yl group, 1H-benzimidazol-4-yl group, benzoxazol-5-yl group,  
 benzoxazol-6-yl group, benzoxazol-4-yl group, benzoxazol-7-yl group, 1H-  
 pyrrolo[3,2-b]pyridin-5-yl group, 1H-pyrrolo[3,2-b]pyridin-6-yl group,  
 benzo[1,2,5]thiadiazol-5-yl group, benzo[1,2,5]thiadiazol-4-yl group, 1H-  
 benzotriazol-5-yl group, 1H-benzotriazol-4-yl group, 1,3-dihydropyrrolo[2,3-  
 b]pyridin-5-yl group, 1,3-dihydropyrrolo[2,3-b]pyridin-4-yl group, 1,3-  
 dihydrobenzimidazol-5-yl group, 1,3-dihydrobenzimidazol-4-yl group, dihydro-3H-  
 benzoxazol-6-yl group, dihydro-3H-benzoxazol-7-yl group, dihydro-3H-benzoxazol-5-  
 yl group, dihydro-3H-benzoxazol-4-yl group, phthalazin-6-yl group, phthalazin-5-yl  
 group, [1,8]naphthalidin-3-yl group, [1,8]naphthalidin-4-yl group,  
 [1,5]naphthalidin-3-yl group, [1,5]naphthalidin-4-yl group, 1H-pyrrolo[3,2-  
 c]pyridin-6-yl group, 1H-pyrrolo[3,2-c]pyridin-4-yl group, 1H-pyrrolo[2,3-c]pyridin-  
 5-yl group, 1H-pyrrolo[2,3-c]pyridin-4-yl group, 1H-pyrazolo[4,3-b]pyridin-5-yl  
 group, 1H-pyrazolo[4,3-b]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-6-yl group,  
 1H-pyrazolo[4,3-c]pyridin-4-yl group, 1H-pyrazolo[3,4-c]pyridin-5-yl group, 1H-  
 pyrazolo[3,4-c]pyridin-4-yl group, 1H-pyrazolo[3,4-b]pyridin-5-yl group, 1H-  
 pyrazolo[3,4-b]pyridin-4-yl group, [1,2,4]triazolo[4,3-a]pyridin-6-yl group,  
 [1,2,4]triazolo[4,3-a]pyridin-7-yl group, thieno[3,2-c]pyridin-2-yl group, thieno[3,2-  
 c]pyridin-3-yl group, thieno[3,2-c]pyridin-6-yl group, thieno[3,2-b]pyridin-2-yl group,  
 thieno[3,2-b]pyridin-3-yl group, thieno[3,2-b]pyridin-5-yl group, thieno[3,2-  
 b]pyridin-6-yl group, 1H-thieno[3,2-c]pyrazol-5-yl group, 1H-thieno[3,2-c]pyrazol-4-  
 yl group, benzo[d]isoxazol-5-yl group, benzo[d]isoxazol-4-yl group, benzo[d]isoxazol-  
 6-yl group, benzo[d]isoxazol-7-yl group, benzo[c]isoxazol-5-yl group,  
 benzo[c]isoxazol-4-yl group, benzo[c]isoxazol-6-yl group, benzo[c]isoxazol-7-yl group,  
 indolizin-7-yl group, indolizin-6-yl group, indolizine-8-yl group, 1,3-dihydroindol-5-  
 yl group, 1,3-dihydroindol-4-yl group, 1,3-dihydroindol-6-yl group, 1H-pyrazolo[3,4-  
 d]thiazol-5-yl group, 2H-isoindol-5-yl group, 2H-isoindol-4-yl group,

[1,2,4]triazolo[1,5-a]pyrimidin-6-yl group, 1H-pyrazolo[3,4-b]pyrazin-5-yl group, 1H-imidazo[4,5-b]pyrazin-5-yl group, 7H-purin-2-yl group, 4H-chromen-6-yl group, or 4H-chromen-5-yl group (these groups may be substituted with one of Xa or two or more of the same or different Xa),

Xa represents oxo group, thioxo group, fluorine atom, chlorine atom, trifluoromethyl group, methyl group, ethyl group, propyl group, 2-hydroxyethyl group, carboxymethyl group, 2-carboxyethyl group, N,N-dimethylcarbamoylmethyl group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, carboxymethyloxy group, 2-carboxyethyloxy group, N,N-dimethylcarbamoylmethyloxy group, amino group, methylamino group, dimethylamino group, 2-hydroxyethylamino group, carbamoylamino group, acetylamino group, furan-2-carboxyamino group, 2-hydroxyacetylamino group, 2-aminoacetylamino group, methylsulfonylamino group, (N,N-dimethylsulfamoyl)amino group, methanesulfonyl group, sulfamoyl group, N-methylsulfamoyl group, N,N-dimethylsulfamoyl group, carboxyl group, acetyl group, carbamoyl group, or N,N-dimethylcarbamoyl group, and

Y is hydrogen atom, methyl group, or ethyl group.

(65) The compound or salt thereof according to (1-2) mentioned above, wherein, in the formula (I), n is an integer of 2,

C<sup>3</sup> is carbon atom to which AR binds, C<sup>4</sup> is carbon atom to which Rs binds, C<sup>2</sup>, C<sup>5</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms,

Rs is -N(Ry)(Rz), Rz is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group,

4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl

group, 2-(N-phenyl-N-methylamino)ethyl group, 2-(N-ethyl-N-phenylamino)ethyl  
 group, isobutyryl group, isopropylthiocarbonyl group, isopropylsulfonyl group,  
 valeryl group, butylthiocarbonyl group, isovaleryl group, isobutylthiocarbonyl group,  
 pivaloyl group, t-butylthiocarbonyl group, cyclopropylcarbonyl group,  
 cyclopropylthiocarbonyl group, cyclopentylcarbonyl group, cyclopentylthiocarbonyl  
 group, cyclohexylcarbonyl group, cyclohexylthiocarbonyl group,  
 cyclopentylmethylcarbonyl group, cyclopentylmethylthiocarbonyl group,  
 cyclohexylmethylcarbonyl group, cyclohexylmethylthiocarbonyl group, benzoyl  
 group, thiobenzoyl group, phenylsulfonyl group, 4-methylphenylcarbonyl group, 4-  
 methylphenylthiocarbonyl group, 4-methylphenylsulfonyl group, 4-  
 chlorophenylcarbonyl group, 4-chlorophenylthiocarbonyl group, 4-  
 fluorophenylcarbonyl group, 4-fluorophenylthiocarbonyl group,  
 isopropoxy carbonyl group, N-isopropylcarbamoyl group, N-isopropylthiocarbamoyl  
 group, butyloxy carbonyl group, N-butylcarbamoyl group, N-butylthiocarbamoyl  
 group, isobutyloxy carbonyl group, N-isobutylcarbamoyl group, N-  
 isobutylthiocarbamoyl group, t-butyloxy carbonyl group, N-t-butylcarbamoyl group,  
 N-t-butylthiocarbamoyl group, cyclopropyloxy carbonyl group, N-  
 cyclopropylcarbamoyl group, N-cyclopropylthiocarbamoyl group,  
 cyclopentyloxy carbonyl group, N-cyclopentylcarbamoyl group, N-  
 cyclopentylthiocarbamoyl group, cyclohexyloxy carbonyl group, N-  
 cyclohexylcarbamoyl group, N-cyclohexylthiocarbamoyl group,  
 cyclopentylmethyloxy carbonyl group, cyclohexylmethyloxy carbonyl group,  
 phenyloxy carbonyl group, N-phenylcarbamoyl group, N-phenylthiocarbamoyl group,  
 4-methylphenyloxy carbonyl group, N-(4-methylphenyl)carbamoyl group, N-(4-  
 methylphenyl)thiocarbamoyl group, 4-chlorophenyloxy carbonyl group, N-(4-  
 chlorophenyl)carbamoyl group, N-(4-chlorophenyl)thiocarbamoyl group, 4-  
 fluorophenyloxy carbonyl group, N-(4-fluorophenyl)carbamoyl group, N-(4-

fluorophenyl)thiocarbamoyl group, (pyrrolidino-1-yl)carbonyl group, (piperidino-1-yl)carbonyl group, or (morpholino-4-yl)carbonyl group, R<sub>y</sub> is hydrogen atom, methyl group, ethyl group or isobutyl group, or binds to R<sub>z</sub> to form pyrrolidino group, piperidino group, or morpholino group together with the nitrogen atom to which they bind,

AR is naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethoxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group, 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-

dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, or benzoxazol-5-yl group, and

Y is hydrogen atom, methyl group, or ethyl group.

(65-2) The compound or salt thereof according to (1-2) mentioned above, wherein, in the formula (I), Link is  $-(CH_2)_n-$ , n is an integer of 2,

C<sup>3</sup> is carbon atom to which AR binds, C<sup>4</sup> is carbon atom to which Rs binds, C<sup>2</sup>, C<sup>5</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms,

Rs is  $-N(R_y)(R_z)$ , and the group represented by  $-N(R_y)(R_z)$  is N,N-dimethylamino group, N-ethyl-N-methylamino group, N,N-diethylamino group, N-methyl-N-propylamino group, N-ethyl-N-propylamino group, N-isopropyl-N-methylamino group, N-ethyl-N-isopropylamino group, N-butylamino group, N-butyl-N-methylamino group, N-butyl-N-ethylamino group, N-isobutylamino group, N-isobutyl-N-methylamino group, N-ethyl-N-isobutylamino group, N-(2-ethylbutyl)amino group, N-(2-ethylbutyl)-N-methylamino group, N-cyclopentylamino group, N-cyclopentyl-N-methylamino group, N-cyclohexylamino group, N-cyclohexyl-N-methylamino group, N-cycloheptylamino group, N-(cyclopentylmethyl)amino group, N-(cyclopentylmethyl)-N-methylamino group, N-(cyclohexylmethyl)amino group, N-(cyclohexylmethyl)-N-methylamino group, N-(2-methylphenyl)amino group, N-(4-methylphenyl)amino group, N-(2-fluorophenyl)amino group, N-(3-fluorophenyl)amino group, N-(4-fluorophenyl)amino

group, N-(2-chlorophenyl)amino group, N-(3-chlorophenyl)amino group, N-(4-chlorophenyl)amino group, N-(indan-2-yl)amino group, N-(1-phenylethyl)amino group, N-[1-(2-fluorophenyl)ethyl]amino group, N-[1-(3-fluorophenyl)ethyl]amino group, N-[1-(4-fluorophenyl)ethyl]amino group, N-[1-(2-chlorophenyl)ethyl]amino group, N-[1-(3-chlorophenyl)ethyl]amino group, N-[1-(4-chlorophenyl)ethyl]amino group, N-(2-methylphenylmethyl)amino group, N-methyl-N-(2-methylphenylmethyl)amino group, N-(3-methylphenylmethyl)amino group, N-methyl-N-(3-methylphenylmethyl)amino group, N-(4-methylphenylmethyl)amino group, N-methyl-N-(4-methylphenylmethyl)amino group, N-(2-fluorophenylmethyl)amino group, N-(2-fluorophenylmethyl)-N-methylamino group, N-(3-fluorophenylmethyl)amino group, N-(3-fluorophenylmethyl)-N-methylamino group, N-(4-fluorophenylmethyl)amino group, N-(4-fluorophenylmethyl)-N-methylamino group, N-(2-chlorophenylmethyl)amino group, N-(2-chlorophenylmethyl)-N-methylamino group, N-(3-chlorophenylmethyl)amino group, N-(3-chlorophenylmethyl)-N-methylamino group, N-(4-chlorophenylmethyl)amino group, N-(4-chlorophenylmethyl)-N-methylamino group, N-(2,3-difluorophenylmethyl)amino group, N-(2,3-difluorophenylmethyl)-N-methylamino group, N-(2,4-difluorophenylmethyl)amino group, N-(2,4-difluorophenylmethyl)-N-methylamino group, N-(2,5-difluorophenylmethyl)amino group, N-(2,5-difluorophenylmethyl)-N-methylamino group, N-(3,4-difluorophenylmethyl)amino group, N-(3,4-difluorophenylmethyl)-N-methylamino group, N-(3,5-difluorophenylmethyl)amino group, N-(3,5-difluorophenylmethyl)-N-methylamino group, N-(2,3-dichlorophenylmethyl)amino group, N-(2,3-dichlorophenylmethyl)-N-methylamino group, N-(2,4-dichlorophenylmethyl)amino group, N-(2,4-dichlorophenylmethyl)-N-methylamino group, N-(2,5-dichlorophenylmethyl)amino group, N-(2,5-dichlorophenylmethyl)-N-methylamino group, N-(2,6-dichlorophenylmethyl)amino group, N-(2,6-dichlorophenylmethyl)-N-methylamino



group, N-(3,4-dichlorophenylmethyl)amino group, N-(3,4-dichlorophenylmethyl)-N-methylamino group, N-(3,5-dichlorophenylmethyl)amino group, N-(3,5-dichlorophenylmethyl)-N-methylamino group, N-[2-(trifluoromethyl)phenylmethyl]amino group, N-methyl-N-[2-(trifluoromethyl)phenylmethyl]amino group, N-[3-(trifluoromethyl)phenylmethyl]amino group, N-methyl-N-[3-(trifluoromethyl)phenylmethyl]amino group, N-[4-(trifluoromethyl)phenylmethyl]amino group, N-methyl-N-[4-(trifluoromethyl)phenylmethyl]amino group, 1-pyrrolidino group, 1-(4-methylpiperidino) group, 1-homopiperidino group, or 4-morpholino group,

AR is naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethoxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-

indol-5-yl group, benzothiazol-6-yl group, 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, or benzoxazol-5-yl group, and

Y is hydrogen atom, methyl group, or ethyl group.

(65-3) The compound or salt thereof according to (1) mentioned above, wherein, in the formula (I), Link is  $-(CH_2)_n-$ , n is an integer of 2,

C<sup>3</sup> is carbon atom to which AR binds, C<sup>4</sup> is carbon atom to which Rs binds, C<sup>5</sup> is nitrogen atom, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms,

Rs is -N(Ry)(Rz), and the group represented by -N(Ry)(Rz) is N,N-dimethylamino group, N-ethyl-N-methylamino group, N,N-diethylamino group, N-methyl-N-propylamino group, N-ethyl-N-propylamino group, N-isopropyl-N-methylamino group, N-ethyl-N-isopropylamino group, N-butylamino group, N-butyl-N-methylamino group, N-butyl-N-ethylamino group, N-isobutylamino group, N-isobutyl-N-methylamino group, N-ethyl-N-isobutylamino group, N-(2-ethylbutyl)amino group, N-(2-ethylbutyl)-N-methylamino group, N-cyclopentylamino group, N-cyclopentyl-N-methylamino group, N-cyclohexylamino

group, N-cyclohexyl-N-methylamino group, N-cycloheptylamino group, N-(cyclopentylmethyl)amino group, N-(cyclopentylmethyl)-N-methylamino group, N-(cyclohexylmethyl)amino group, N-(cyclohexylmethyl)-N-methylamino group, N-(2-methylphenyl)amino group, N-(4-methylphenyl)amino group, N-(2-fluorophenyl)amino group, N-(3-fluorophenyl)amino group, N-(4-fluorophenyl)amino group, N-(2-chlorophenyl)amino group, N-(3-chlorophenyl)amino group, N-(4-chlorophenyl)amino group, N-(indan-2-yl)amino group, N-(1-phenylethyl)amino group, N-[1-(2-fluorophenyl)ethyl]amino group, N-[1-(3-fluorophenyl)ethyl]amino group, N-[1-(4-fluorophenyl)ethyl]amino group, N-[1-(2-chlorophenyl)ethyl]amino group, N-[1-(3-chlorophenyl)ethyl]amino group, N-[1-(4-chlorophenyl)ethyl]amino group, N-(2-methylphenylmethyl)amino group, N-methyl-N-(2-methylphenylmethyl)amino group, N-(3-methylphenylmethyl)amino group, N-methyl-N-(3-methylphenylmethyl)amino group, N-(4-methylphenylmethyl)amino group, N-methyl-N-(4-methylphenylmethyl)amino group, N-(2-fluorophenylmethyl)amino group, N-(2-fluorophenylmethyl)-N-methylamino group, N-(3-fluorophenylmethyl)amino group, N-(3-fluorophenylmethyl)-N-methylamino group, N-(4-fluorophenylmethyl)amino group, N-(4-fluorophenylmethyl)-N-methylamino group, N-(2-chlorophenylmethyl)amino group, N-(2-chlorophenylmethyl)-N-methylamino group, N-(3-chlorophenylmethyl)amino group, N-(3-chlorophenylmethyl)-N-methylamino group, N-(4-chlorophenylmethyl)amino group, N-(4-chlorophenylmethyl)-N-methylamino group, N-(2,3-difluorophenylmethyl)amino group, N-(2,3-difluorophenylmethyl)-N-methylamino group, N-(2,4-difluorophenylmethyl)amino group, N-(2,4-difluorophenylmethyl)-N-methylamino group, N-(2,5-difluorophenylmethyl)amino group, N-(2,5-difluorophenylmethyl)-N-methylamino group, N-(3,4-difluorophenylmethyl)amino group, N-(3,4-difluorophenylmethyl)-N-methylamino group, N-(3,5-difluorophenylmethyl)amino group, N-(3,5-difluorophenylmethyl)-N-methylamino

group, N-(2,3-dichlorophenylmethyl)amino group, N-(2,3-dichlorophenylmethyl)-N-methylamino group, N-(2,4-dichlorophenylmethyl)amino group, N-(2,4-dichlorophenylmethyl)-N-methylamino group, N-(2,5-dichlorophenylmethyl)amino group, N-(2,5-dichlorophenylmethyl)-N-methylamino group, N-(2,6-dichlorophenylmethyl)amino group, N-(2,6-dichlorophenylmethyl)-N-methylamino group, N-(3,4-dichlorophenylmethyl)amino group, N-(3,4-dichlorophenylmethyl)-N-methylamino group, N-(3,5-dichlorophenylmethyl)amino group, N-(3,5-dichlorophenylmethyl)-N-methylamino group, N-[2-(trifluoromethyl)phenylmethyl]amino group, N-methyl-N-[2-(trifluoromethyl)phenylmethyl]amino group, N-[3-(trifluoromethyl)phenylmethyl]amino group, N-methyl-N-[3-(trifluoromethyl)phenylmethyl]amino group, N-[4-(trifluoromethyl)phenylmethyl]amino group, N-methyl-N-[4-(trifluoromethyl)phenylmethyl]amino group, 1-pyrrolidino group, 1-(4-methylpiperidino) group, 1-homopiperidino group, or 4-morpholino group,

AR is naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-

methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group, 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, or benzoxazol-5-yl group, and

Y is hydrogen atom, methyl group, or ethyl group.

(66) The compound or salt thereof according to (1-2) mentioned above, wherein, in the formula (I), n is an integer of 2, AR binds to C<sup>3</sup> in the aromatic ring (E), R<sub>s</sub> binds to C<sup>4</sup> in the aromatic ring (E), C<sup>5</sup> is carbon atom substituted with N(R<sub>n</sub><sup>1</sup>)(R<sub>n</sub><sup>2</sup>) (provided that one of R<sub>n</sub><sup>1</sup> and R<sub>n</sub><sup>2</sup> is a substituent other than hydrogen atom), C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms, R<sub>s</sub> is -O-R<sub>x</sub>, and Y is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms.

(67) The compound or salt thereof according to (4) mentioned above, wherein, in the formula (I), Link is  $-(CH_2)_n-$ ,  $n$  is an integer of 2, AR binds to  $C^3$  in the aromatic ring (E), Rs binds to  $C^4$  in the aromatic ring (E),  $C^5$  is carbon atom substituted with  $-N(Rn^1)(Rn^2)$  (provided that one of  $Rn^1$  and  $Rn^2$  is a substituent other than hydrogen atom),  $C^2$  and  $C^6$  are unsubstituted ring-constituting carbon atoms, Rs is  $-O-Rx$ , and Y is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms.

(68) The compound or salt thereof according to (5) mentioned above, wherein, in the formula (I), Link is  $-(CH_2)_n-$ ,  $n$  is an integer of 2, AR binds to  $C^3$  in the aromatic ring (E), Rs binds to  $C^4$  in the aromatic ring (E),  $C^5$  is carbon atom substituted with  $-N(Rn^1)(Rn^2)$  (provided that one of  $Rn^1$  and  $Rn^2$  is a substituent other than hydrogen atom),  $C^2$  and  $C^6$  are unsubstituted ring-constituting carbon atoms, Rs is  $-O-Rx$ , and Y is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms.

(69) The compound or salt thereof according to any one of (66) to (68) mentioned above, wherein, in the formula (I), Xa which may substitute on AR is methyl group, ethyl group, propyl group, hydroxyethyl group, carboxymethyl group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, amino group, methylamino group, dimethylamino group, carboxyl group, carbamoyl group, acetyl group, methanesulfonyl group, sulfamoyl group, or N,N-dimethylsulfamoyl group.

(70) The compound or salt thereof according to any one of (66) to (69) mentioned above, wherein, in the formula (I), Rs is  $-O-Rx$ , Rx is a group selected from butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, and cyclohexylmethyl group, or Rb (provided that Q in Rb is phenyl group or indan-2-yl group),  $A^1$  is a single bond, or methylene group substituted with methyl group or ethyl group or unsubstituted methylene group, or ethylene group substituted with methyl group or ethyl group or unsubstituted ethylene group,  $A^2$  represents a single bond, oxygen atom, sulfur atom,  $-N(methyl)-$ , or  $-N(ethyl)-$  (provided that when  $A^2$  is oxygen atom, sulfur atom,

-N(methyl)-, or -N(ethyl)-, A<sup>1</sup> is ethylene), and R<sup>2</sup> and R<sup>3</sup> independently represent hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, or dimethylamino group (provided that when Q is phenyl group, A<sup>1</sup> is a single bond or unsubstituted methylene, and A<sup>2</sup> is a single bond, one of R<sup>2</sup> and R<sup>3</sup> is a substituent other than hydrogen atom).

(71) The compound or salt thereof according to (1-2) mentioned above, wherein, in the formula (I), n is an integer of 2,

C<sup>3</sup> is carbon atom to which AR binds, C<sup>4</sup> is carbon atom to which Rs binds, C<sup>5</sup> is carbon atom substituted with Zx, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms,

Zx is N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, or N,N-dimethylsulfamoylamino group,

Rs is -O-Rx, Rx is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl

group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, or 2-(N-ethyl-N-phenylamino)ethyl group,

AR is naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl



group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group, 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, or benzoxazol-5-yl group, and

Y is hydrogen atom, methyl group, or ethyl group.

(72) The compound or salt thereof according to (1-2) mentioned above, wherein, in the formula (I), n is an integer of 2, AR binds to C<sup>3</sup> in the aromatic ring (E), Rs binds to C<sup>4</sup> in the aromatic ring (E), C<sup>5</sup> is a ring-constituting carbon atom substituted with Zx, or an unsubstituted ring-constituting carbon atom, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms, Rs is -D-Rc, D is oxygen atom or sulfur atom, and Y is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms.

(73) The compound or salt thereof according to (4) mentioned above, wherein, in the formula (I), Link is -(CH<sub>2</sub>)<sub>n</sub>-, n is an integer of 2, AR binds to C<sup>3</sup> in the aromatic ring (E), Rs binds to C<sup>4</sup> in the aromatic ring (E), C<sup>5</sup> is a ring-constituting carbon atom substituted with Zx, or an unsubstituted ring-constituting carbon atom, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms, Rs is -O-Rc, and Y is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms.

(74) The compound or salt thereof according to (5) mentioned above, wherein, in the formula (I), Link is -(CH<sub>2</sub>)<sub>n</sub>-, n is an integer of 2, AR binds to C<sup>3</sup> in the aromatic ring (E), Rs binds to C<sup>4</sup> in the aromatic ring (E), C<sup>5</sup> is a ring-constituting carbon atom substituted with Zx, or an unsubstituted ring-constituting carbon atom, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms, Rs is -O-Rc, and Y is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms.

(75) The compound or salt thereof according to any one of (72) to (74) mentioned above, wherein, in the formula (I), Xa which may substitute on AR is methyl group, ethyl group, propyl group, hydroxyethyl group, carboxymethyl group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, amino group, methylamino group, dimethylamino group, carboxyl group, carbamoyl group, acetyl group, methanesulfonyl group, sulfamoyl group, or N,N-dimethylsulfamoyl group.

(76) The compound or salt thereof according to (1-2) mentioned above, wherein, in

the formula (I), n is an integer of 1 to 3,

C<sup>3</sup> is carbon atom to which AR binds, C<sup>4</sup> is carbon atom to which Rs binds, C<sup>5</sup> may be replaced with V, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms,

V is nitrogen atom, or carbon atom substituted with Z<sub>x</sub>, Z<sub>x</sub> is fluorine atom, chlorine atom, bromine atom, nitro group, methyl group, hydroxyl group, methoxy group, amino group, N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, or N,N-dimethylsulfamoylamino group,

Rs is -D-R<sub>c</sub>, D is oxygen atom or sulfur atom, p in R<sub>c</sub> is an integer of 2 or 3, A<sup>4</sup> is a single bond or methylene, A<sup>5</sup> is -C(O)-, -C(S)-, or -S(O)<sub>2</sub>-, R<sub>d</sub> is hydrogen atom, or methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group, cyclopropylmethyl group, cyclopentyl group, cyclopentylmethyl group, cyclohexyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, or pyridin-4-yl group, R<sub>e</sub> is methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, phenylmethyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, furan-2-yl group, furan-3-yl group, thiophen-2-yl group, thiophen-3-yl group, methoxy group, ethoxy group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenyloxy group, 4-

methylphenoxy group, 4-chlorophenoxy group, 4-fluorophenoxy group,  
 thiomethoxy group, amino group, N-methylamino group, N,N-dimethylamino group,  
 N-ethylamino group, N,N-diethylamino group, N-propylamino group, N-  
 isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino  
 group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino  
 group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-  
 chlorophenyl)amino group, N-(4-fluorophenyl)amino group, N-(pyridin-2-yl)amino  
 group, N-(pyridin-3-yl)amino group, N-(pyridin-4-yl)amino group, N-(furan-2-  
 yl)amino group, N-(furan-3-yl)amino group, N-(thiophen-2-yl)amino group, N-  
 (thiophen-3-yl)amino group, pyrrolidino group, piperidino group, morpholino group,  
 methyloxycarbonylamino group, or ethyloxycarbonylamino group,

AR is naphthalen-2-yl group, naphthalen-1-yl group, benzofuran-5-yl group,  
 benzofuran-4-yl group, benzofuran-2-yl group, benzo[b]thiophen-5-yl group,  
 benzo[b]thiophen-4-yl group, benzo[b]thiophen-2-yl group, indol-5-yl group, indol-4-  
 yl group, indol-6-yl group, benzothiazol-6-yl group, benzothiazol-7-yl group,  
 benzothiazol-5-yl group, benzothiazol-4-yl group, dihydro-3H-benzothiazol-6-yl  
 group, dihydro-3H-benzothiazol-7-yl group, dihydro-3H-benzothiazol-5-yl group,  
 dihydro-3H-benzothiazol-4-yl group, quinolin-6-yl group, quinolin-3-yl group,  
 quinolin-5-yl group, quinolin-7-yl group, dihydro-1H-quinolin-6-yl group, dihydro-  
 1H-quinolin-5-yl group, benzo[d]isothiazol-5-yl group, benzo[d]isothiazol-4-yl group,  
 benzo[d]isothiazol-6-yl group, benzo[d]isothiazol-7-yl group, 1H-indazol-5-yl group,  
 1H-indazol-4-yl group, 1H-indazol-6-yl group, benzo[c]isothiazol-5-yl group,  
 benzo[c]isothiazol-4-yl group, benzo[c]isothiazol-6-yl group, benzo[c]isothiazol-7-yl  
 group, 2H-indazol-5-yl group, 2H-indazol-4-yl group, 2H-indazol-6-yl group,  
 imidazo[1,2-a]pyridin-6-yl group, imidazo[1,2-a]pyridin-7-yl group, 1H-pyrrolo[2,3-  
 b]pyridin-5-yl group, 1H-pyrrolo[2,3-b]pyridin-4-yl group, isoquinolin-6-yl group,  
 isoquinolin-3-yl group, isoquinolin-5-yl group, isoquinolin-7-yl group, dihydro-2H-

isoquinolin-6-yl group, dihydro-2H-isoquinolin-5-yl group, cinnolin-6-yl group,  
 cinnolin-5-yl group, quinazolin-6-yl group, quinazolin-7-yl group, quinazolin-5-yl  
 group, quinoxalin-2-yl group, quinoxalin-6-yl group, quinoxalin-5-yl group, 1H-  
 benzimidazol-5-yl group, 1H-benzimidazol-4-yl group, benzoxazol-5-yl group,  
 benzoxazol-6-yl group, benzoxazol-4-yl group, benzoxazol-7-yl group, 1H-  
 pyrrolo[3,2-b]pyridin-5-yl group, 1H-pyrrolo[3,2-b]pyridin-6-yl group,  
 benzo[1,2,5]thiadiazol-5-yl group, benzo[1,2,5]thiadiazol-4-yl group, 1H-  
 benzotriazol-5-yl group, 1H-benzotriazol-4-yl group, 1,3-dihydropyrrolo[2,3-  
 b]pyridin-5-yl group, 1,3-dihydropyrrolo[2,3-b]pyridin-4-yl group, 1,3-  
 dihydrobenzimidazol-5-yl group, 1,3-dihydrobenzimidazol-4-yl group, dihydro-3H-  
 benzoxazol-6-yl group, dihydro-3H-benzoxazol-7-yl group, dihydro-3H-benzoxazol-5-  
 yl group, dihydro-3H-benzoxazol-4-yl group, phthalazin-6-yl group, phthalazin-5-yl  
 group, [1,8]naphthalidin-3-yl group, [1,8]naphthalidin-4-yl group,  
 [1,5]naphthalidin-3-yl group, [1,5]naphthalidin-4-yl group, 1H-pyrrolo[3,2-  
 c]pyridin-6-yl group, 1H-pyrrolo[3,2-c]pyridin-4-yl group, 1H-pyrrolo[2,3-c]pyridin-  
 5-yl group, 1H-pyrrolo[2,3-c]pyridin-4-yl group, 1H-pyrazolo[4,3-b]pyridin-5-yl  
 group, 1H-pyrazolo[4,3-b]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-6-yl group,  
 1H-pyrazolo[4,3-c]pyridin-4-yl group, 1H-pyrazolo[3,4-c]pyridin-5-yl group, 1H-  
 pyrazolo[3,4-c]pyridin-4-yl group, 1H-pyrazolo[3,4-b]pyridin-5-yl group, 1H-  
 pyrazolo[3,4-b]pyridin-4-yl group, [1,2,4]triazolo[4,3-a]pyridin-6-yl group,  
 [1,2,4]triazolo[4,3-a]pyridin-7-yl group, thieno[3,2-c]pyridin-2-yl group, thieno[3,2-  
 c]pyridin-3-yl group, thieno[3,2-c]pyridin-6-yl group, thieno[3,2-b]pyridin-2-yl group,  
 thieno[3,2-b]pyridin-3-yl group, thieno[3,2-b]pyridin-5-yl group, thieno[3,2-  
 b]pyridin-6-yl group, 1H-thieno[3,2-c]pyrazol-5-yl group, 1H-thieno[3,2-c]pyrazol-4-  
 yl group, benzo[d]isoxazol-5-yl group, benzo[d]isoxazol-4-yl group, benzo[d]isoxazol-  
 6-yl group, benzo[d]isoxazol-7-yl group, benzo[c]isoxazol-5-yl group,  
 benzo[c]isoxazol-4-yl group, benzo[c]isoxazol-6-yl group, benzo[c]isoxazol-7-yl group,

indolizin-7-yl group, indolizin-6-yl group, indolizine-8-yl group, 1,3-dihydroindol-5-yl group, 1,3-dihydroindol-4-yl group, 1,3-dihydroindol-6-yl group, 1H-pyrazolo[3,4-d]thiazol-5-yl group, 2H-isoindol-5-yl group, 2H-isoindol-4-yl group, [1,2,4]triazolo[1,5-a]pyrimidin-6-yl group, 1H-pyrazolo[3,4-b]pyrazin-5-yl group, 1H-imidazo[4,5-b]pyrazin-5-yl group, 7H-purin-2-yl group, 4H-chromen-6-yl group, or 4H-chromen-5-yl group (the aforementioned groups may be substituted with one of Xa or two or more of the same or different Xa), Xa is oxo group, thioxo group, fluorine atom, chlorine atom, trifluoromethyl group, methyl group, ethyl group, propyl group, 2-hydroxyethyl group, carboxymethyl group, 2-carboxyethyl group, N,N-dimethylcarbamoylmethyl group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, carboxymethyloxy group, 2-carboxyethyloxy group, N,N-dimethylcarbamoylmethyloxy group, amino group, methylamino group, dimethylamino group, 2-hydroxyethylamino group, carbamoylamino group, acetylamino group, furan-2-carboxyamino group, 2-hydroxyacetylamino group, 2-aminoacetylamino group, methylsulfonylamino group, (N,N-dimethylsulfamoyl)amino group, methanesulfonyl group, sulfamoyl group, N-methylsulfamoyl group, N,N-dimethylsulfamoyl group, carboxyl group, acetyl group, carbamoyl group, or N,N-dimethylcarbamoyl group, and

Y is hydrogen atom, methyl group, or ethyl group.

(77) The compound or salt thereof according to (1-2) mentioned above, wherein, in the formula (I), n is an integer of 2,

C<sup>3</sup> is carbon atom to which AR binds, C<sup>4</sup> is carbon atom to which Rs binds, C<sup>5</sup> may be replaced with V, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms,

V is nitrogen atom, or carbon atom substituted with Zx, Zx is fluorine atom, chlorine atom, bromine atom, methyl group, hydroxyl group, methoxy group, amino group, N-methylamino group, N-ethylamino group, N-propylamino group, N-

isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, or N,N-dimethylsulfamoylamino group,

Rs is -O-Rc, p in Rc is an integer of 2, A<sup>4</sup> is a single bond or methylene, A<sup>5</sup> is -C(O)-, -C(S)-, or -S(O)<sub>2</sub>-, Rd is methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, or 4-fluorophenylmethyl group, Re is isopropyl group, butyl group, isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenyloxy group, 4-methylphenyloxy group, 4-chlorophenyloxy group, 4-fluorophenyloxy group, N-propylamino group, N-isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group, N-(4-fluorophenyl)amino group, pyrrolidino group, piperidino group, or morpholino group,

AR is naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-

methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group, 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, or benzoxazol-5-yl group, and

Y is hydrogen atom, methyl group, or ethyl group.

(78) The compound or salt thereof according to (7) mentioned above, wherein, in the formula (I), Link is  $-(CH_2)_n-$ , n is an integer of 2, AR binds to C<sup>3</sup> in the aromatic



ring (E), Rs binds to C<sup>4</sup> in the aromatic ring (E), C<sup>5</sup> is a ring-constituting carbon atom substituted with Zx, or an unsubstituted ring-constituting carbon atom, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms, Rs is -O-Rx, and Y is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms.

(79) The compound or salt thereof according to (78) mentioned above, wherein, in the formula (I), Xa which may substitute on AR is methyl group, ethyl group, propyl group, hydroxyethyl group, carboxymethyl group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, amino group, methylamino group, dimethylamino group, carboxyl group, carbamoyl group, acetyl group, methanesulfonyl group, sulfamoyl group, or N,N-dimethylsulfamoyl group.

(80) The compound or salt thereof according to (78) or (79) mentioned above, wherein, in the formula (I), Rs is -O-Rx, Rx is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, or cyclohexylmethyl group, or Rb (provided that Q in Rb is phenyl group or indan-2-yl group), A<sup>1</sup> is a single bond, or methylene group substituted with methyl group or ethyl group or unsubstituted methylene group, or ethylene group substituted with methyl group or ethyl group or unsubstituted ethylene group, A<sup>2</sup> is a single bond, oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)- (provided that when A<sup>2</sup> is oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)-, A<sup>1</sup> is ethylene), and R<sup>2</sup> and R<sup>3</sup> independently represent hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, or dimethylamino group (provided that when Q is phenyl group, A<sup>1</sup> is a single bond or unsubstituted methylene, and A<sup>2</sup> is a single bond, one of R<sup>2</sup> and R<sup>3</sup> is a substituent other than hydrogen atom).

(81) The compound or salt thereof according to (1-2) mentioned above, wherein, in the formula (I), n is an integer of 1 to 3,

C<sup>3</sup> is carbon atom to which AR binds, C<sup>4</sup> is carbon atom to which Rs binds,

C<sup>5</sup> may be replaced with V, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms,

V is nitrogen atom, or carbon atom substituted with Zx, Zx is fluorine atom, chlorine atom, bromine atom, nitro group, methyl group, hydroxyl group, methoxy group, amino group, N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, or N,N-dimethylsulfamoylamino group,

Rs is -O-Rx, Rx is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-cyclopentylethyl group, or 2-cyclohexylethyl group, or Rb or Rc, Q in Rb is phenyl group, thienyl group, furyl group, pyridyl group, oxazolyl group, naphthyl group, tetrahydronaphthyl group, indanyl group, indolyl group, or dihydrobenzodioxyl group, A<sup>2</sup> is a single bond, oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)- (provided that when A<sup>2</sup> is oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)-, A<sup>1</sup> is ethylene), R<sup>2</sup> and R<sup>3</sup> independently represent hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, dimethylamino group, acetylamino group, or methylsulfonylamino group (provided that when Q is phenyl group, A<sup>1</sup> is a single bond or unsubstituted methylene, and A<sup>2</sup> is a single bond, one of R<sup>2</sup> and R<sup>3</sup> is a substituent other than hydrogen atom), p in Rc is an integer of 2 or 3, A<sup>4</sup> is a single bond or methylene, A<sup>5</sup> is -C(O)-, -C(S)-, or -S(O)<sub>2</sub>-, Rd is hydrogen atom, or methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group, cyclopropylmethyl group, cyclopentyl group, cyclopentylmethyl group, cyclohexyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, or

pyridin-4-yl group, Re is methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, phenylmethyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, furan-2-yl group, furan-3-yl group, thiophen-2-yl group, thiophen-3-yl group, methoxy group, ethoxy group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenyloxy group, 4-methylphenyloxy group, 4-chlorophenyloxy group, 4-fluorophenyloxy group, thiomethoxy group, amino group, N-methylamino group, N,N-dimethylamino group, N-ethylamino group, N,N-diethylamino group, N-propylamino group, N-isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group, N-(4-fluorophenyl)amino group, N-(pyridin-2-yl)amino group, N-(pyridin-3-yl)amino group, N-(pyridin-4-yl)amino group, N-(furan-2-yl)amino group, N-(furan-3-yl)amino group, N-(thiophen-2-yl)amino group, N-(thiophen-3-yl)amino group, pyrrolidino group, piperidino group, morpholino group, methyloxycarbonylamino group, or ethyloxycarbonylamino group,

AR is cinnolin-6-yl group, cinnolin-5-yl group, quinazolin-6-yl group, quinazolin-7-yl group, quinazolin-5-yl group, quinoxalin-2-yl group, quinoxalin-6-yl group, quinoxalin-5-yl group, 1H-benzimidazol-5-yl group, 1H-benzimidazol-4-yl group, benzoxazol-5-yl group, benzoxazol-6-yl group, benzoxazol-4-yl group, benzoxazol-7-yl group, 1H-pyrrolo[3,2-b]pyridin-5-yl group, 1H-pyrrolo[3,2-b]pyridin-6-yl group, benzo[1,2,5]thiadiazol-5-yl group, benzo[1,2,5]thiadiazol-4-yl

group, 1H-benzotriazol-5-yl group, 1H-benzotriazol-4-yl group, 1,3-dihydropyrrolo[2,3-b]pyridin-5-yl group, 1,3-dihydropyrrolo[2,3-b]pyridin-4-yl group, 1,3-dihydrobenzimidazol-5-yl group, 1,3-dihydrobenzimidazol-4-yl group, dihydro-3H-benzoxazol-6-yl group, dihydro-3H-benzoxazol-7-yl group, dihydro-3H-benzoxazol-5-yl group, dihydro-3H-benzoxazol-4-yl group, phthalazin-6-yl group, phthalazin-5-yl group, [1,8]naphthalidin-3-yl group, [1,8]naphthalidin-4-yl group, [1,5]naphthalidin-3-yl group, [1,5]naphthalidin-4-yl group, 1H-pyrrolo[3,2-c]pyridin-6-yl group, 1H-pyrrolo[3,2-c]pyridin-4-yl group, 1H-pyrrolo[2,3-c]pyridin-5-yl group, 1H-pyrrolo[2,3-c]pyridin-4-yl group, 1H-pyrazolo[4,3-b]pyridin-5-yl group, 1H-pyrazolo[4,3-b]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-4-yl group, 1H-pyrazolo[3,4-c]pyridin-5-yl group, 1H-pyrazolo[3,4-c]pyridin-4-yl group, 1H-pyrazolo[3,4-b]pyridin-5-yl group, 1H-pyrazolo[3,4-b]pyridin-4-yl group, [1,2,4]triazolo[4,3-a]pyridin-6-yl group, [1,2,4]triazolo[4,3-a]pyridin-7-yl group, thieno[3,2-c]pyridin-2-yl group, thieno[3,2-c]pyridin-3-yl group, thieno[3,2-c]pyridin-6-yl group, thieno[3,2-b]pyridin-2-yl group, thieno[3,2-b]pyridin-3-yl group, thieno[3,2-b]pyridin-5-yl group, thieno[3,2-b]pyridin-6-yl group, 1H-thieno[3,2-c]pyrazol-5-yl group, 1H-thieno[3,2-c]pyrazol-4-yl group, benzo[d]isoxazol-5-yl group, benzo[d]isoxazol-4-yl group, benzo[d]isoxazol-6-yl group, benzo[d]isoxazol-7-yl group, benzo[c]isoxazol-5-yl group, benzo[c]isoxazol-4-yl group, benzo[c]isoxazol-6-yl group, benzo[c]isoxazol-7-yl group, indolizin-7-yl group, indolizin-6-yl group, indolizine-8-yl group, 1,3-dihydroindol-5-yl group, 1,3-dihydroindol-4-yl group, 1,3-dihydroindol-6-yl group, 1H-pyrazolo[3,4-d]thiazol-5-yl group, 2H-isoindol-5-yl group, 2H-isoindol-4-yl group, [1,2,4]triazolo[1,5-a]pyrimidin-6-yl group, 1H-pyrazolo[3,4-b]pyrazin-5-yl group, 1H-imidazo[4,5-b]pyrazin-5-yl group, 7H-purin-2-yl group, 4H-chromen-6-yl group, or 4H-chromen-5-yl group (the aforementioned groups may be substituted with one of Xa or two or more of the same or different Xa), Xa is oxo group, thioxo group,

fluorine atom, chlorine atom, trifluoromethyl group, methyl group, ethyl group, propyl group, 2-hydroxyethyl group, carboxymethyl group, 2-carboxyethyl group, N,N-dimethylcarbamoylmethyl group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, carboxymethyloxy group, 2-carboxyethyloxy group, N,N-dimethylcarbamoylmethyloxy group, amino group, methylamino group, dimethylamino group, 2-hydroxyethylamino group, carbamoylamino group, acetylamino group, furan-2-carboxyamino group, 2-hydroxyacetylamino group, 2-aminoacetylamino group, methylsulfonylamino group, (N,N-dimethylsulfamoyl)amino group, methanesulfonyl group, sulfamoyl group, N-methylsulfamoyl group, N,N-dimethylsulfamoyl group, carboxyl group, acetyl group, carbamoyl group, or N,N-dimethylcarbamoyl group, and

Y is hydrogen atom, methyl group, or ethyl group.

(82) The compound or salt thereof according to (6) mentioned above, wherein, in the formula (I), Link is  $-(CH_2)_n-$ , AR binds to C<sup>3</sup> in the aromatic ring (E), Rs binds to C<sup>4</sup> in the aromatic ring (E), C<sup>5</sup> is a ring-constituting carbon atom substituted with Zx, or an unsubstituted ring-constituting carbon atom, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms, Rs is  $-O-R_x$ , and R<sub>x</sub> is a linear or branched saturated alkyl group having 3 to 8 carbon atoms, or Ra or Rb.

(83) The compound or salt thereof according to (82) mentioned above, wherein, in the formula (I), Zx is fluorine atom, chlorine atom, nitro group, amino group, methyl group, or OR<sup>9</sup>.

(84) The compound or salt thereof according to (1-2) mentioned above, wherein, in the formula (I), n is an integer of 1 to 3, AR binds to C<sup>3</sup>, Rs binds to one of the ring-constituting atoms C<sup>4</sup>, C<sup>5</sup>, and C<sup>6</sup>, a ring-constituting carbon atom to which Rs does not bind among C<sup>4</sup>, C<sup>5</sup>, and C<sup>6</sup> may be replaced with V,

V is nitrogen atom or carbon atom substituted with Zx, Zx is fluorine atom, chlorine atom, bromine atom, nitro group, methyl group, hydroxyl group, methoxy

group, amino group, N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, or N,N-dimethylsulfamoylamino group,

Rs is -D-Rx or -N(Ry)(Rz), D is oxygen atom or sulfur atom, Rx is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-cyclopentylethyl group, or 2-cyclohexylethyl group, or Rb or Rc, Q in Rb is phenyl group, thienyl group, furyl group, pyridyl group, oxazolyl group, naphthyl group, tetrahydronaphthyl group, indanyl group, indolyl group, or dihydrobenzodioxyl group, A<sup>2</sup> is a single bind, oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)- (provided that when A<sup>2</sup> is oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)-, A<sup>1</sup> is ethylene), R<sup>2</sup> and R<sup>3</sup> independently represent hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, dimethylamino group, acetylamino group, or methylsulfonylamino group (provided that when Q is phenyl group, A<sup>1</sup> is a single bind or unsubstituted methylene, and A<sup>2</sup> is a single bind, one of R<sup>2</sup> and R<sup>3</sup> is a substituent other than hydrogen atom), p in Rc is an integer of 2 or 3, A<sup>4</sup> is a single bind or methylene, A<sup>5</sup> is -C(O)-, -C(S)-, or -S(O)<sub>2</sub>-, Rd is hydrogen atom, or methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group, cyclopropylmethyl group, cyclopentyl group, cyclopentylmethyl group, cyclohexyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, or pyridin-4-yl group, Re is methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-

chlorophenyl group, 4-fluorophenyl group, phenylmethyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, furan-2-yl group, furan-3-yl group, thiophen-2-yl group, thiophen-3-yl group, methoxy group, ethoxy group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenyloxy group, 4-methylphenyloxy group, 4-chlorophenyloxy group, 4-fluorophenyloxy group, thiomethoxy group, amino group, N-methylamino group, N,N-dimethylamino group, N-ethylamino group, N,N-diethylamino group, N-propylamino group, N-isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group, N-(4-fluorophenyl)amino group, N-(pyridin-2-yl)amino group, N-(pyridin-3-yl)amino group, N-(pyridin-4-yl)amino group, N-(furan-2-yl)amino group, N-(furan-3-yl)amino group, N-(thiophen-2-yl)amino group, N-(thiophen-3-yl)amino group, pyrrolidino group, piperidino group, morpholino group, methyloxycarbonylamino group or ethyloxycarbonylamino group, R<sub>z</sub> is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-

dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, 2-(N-ethyl-N-phenylamino)ethyl group, isobutyryl group, isopropylthiocarbonyl group, isopropylsulfonyl group, valeryl group, butylthiocarbonyl group, isovaleryl group, isobutylthiocarbonyl group,



pivaloyl group, t-butylthiocarbonyl group, cyclopropylcarbonyl group,  
 cyclopropylthiocarbonyl group, cyclopentylcarbonyl group, cyclopentylthiocarbonyl  
 group, cyclohexylcarbonyl group, cyclohexylthiocarbonyl group,  
 cyclopentylmethylcarbonyl group, cyclopentylmethylthiocarbonyl group,  
 cyclohexylmethylcarbonyl group, cyclohexylmethylthiocarbonyl group, benzoyl  
 group, thiobenzoyl group, phenylsulfonyl group, 4-methylphenylcarbonyl group, 4-  
 methylphenylthiocarbonyl group, 4-methylphenylsulfonyl group, 4-  
 chlorophenylcarbonyl group, 4-chlorophenylthiocarbonyl group, 4-  
 fluorophenylcarbonyl group, 4-fluorophenylthiocarbonyl group,  
 isopropoxy carbonyl group, N-isopropylcarbamoyl group, N-isopropylthiocarbamoyl  
 group, butyloxy carbonyl group, N-butylcarbamoyl group, N-butylthiocarbamoyl  
 group, isobutyloxy carbonyl group, N-isobutylcarbamoyl group, N-  
 isobutylthiocarbamoyl group, t-butyloxy carbonyl group, N-t-butylcarbamoyl group,  
 N-t-butylthiocarbamoyl group, cyclopropyloxy carbonyl group, N-  
 cyclopropylcarbamoyl group, N-cyclopropylthiocarbamoyl group,  
 cyclopentyloxy carbonyl group, N-cyclopentylcarbamoyl group, N-  
 cyclopentylthiocarbamoyl group, cyclohexyloxy carbonyl group, N-  
 cyclohexylcarbamoyl group, N-cyclohexylthiocarbamoyl group,  
 cyclopentylmethyloxy carbonyl group, cyclohexylmethyloxy carbonyl group,  
 phenyloxy carbonyl group, N-phenylcarbamoyl group, N-phenylthiocarbamoyl group,  
 4-methylphenyloxy carbonyl group, N-(4-methylphenyl)carbamoyl group, N-(4-  
 methylphenyl)thiocarbamoyl group, 4-chlorophenyloxy carbonyl group, N-(4-  
 chlorophenyl)carbamoyl group, N-(4-chlorophenyl)thiocarbamoyl group, 4-  
 fluorophenyloxy carbonyl group, N-(4-fluorophenyl)carbamoyl group, N-(4-  
 fluorophenyl)thiocarbamoyl group, (pyrrolidino-1-yl)carbonyl group, (piperidino-1-  
 yl)carbonyl group, or (morpholino-4-yl)carbonyl group, Ry is hydrogen atom, methyl  
 group, ethyl group or isobutyl group, or binds to Rz to form pyrrolidino group,

piperidino group, piperazino group, morpholino group, pyrrol-1-yl group, imidazol-1-yl group, or pyrazol-1-yl group together with the nitrogen atom,

AR is naphthalen-2-yl group, naphthalen-1-yl group, benzofuran-5-yl group, benzofuran-4-yl group, benzofuran-2-yl group, benzo[b]thiophen-5-yl group, benzo[b]thiophen-4-yl group, benzo[b]thiophen-2-yl group, indol-5-yl group, indol-4-yl group, indol-6-yl group, benzothiazol-6-yl group, benzothiazol-7-yl group, benzothiazol-5-yl group, benzothiazol-4-yl group, dihydro-3H-benzothiazol-6-yl group, dihydro-3H-benzothiazol-7-yl group, dihydro-3H-benzothiazol-5-yl group, dihydro-3H-benzothiazol-4-yl group, quinolin-6-yl group, quinolin-3-yl group, quinolin-5-yl group, quinolin-7-yl group, dihydro-1H-quinolin-6-yl group, dihydro-1H-quinolin-5-yl group, benzo[d]isothiazol-5-yl group, benzo[d]isothiazol-4-yl group, benzo[d]isothiazol-6-yl group, benzo[d]isothiazol-7-yl group, 1H-indazol-5-yl group, 1H-indazol-4-yl group, 1H-indazol-6-yl group, benzo[c]isothiazol-5-yl group, benzo[c]isothiazol-4-yl group, benzo[c]isothiazol-6-yl group, benzo[c]isothiazol-7-yl group, 2H-indazol-5-yl group, 2H-indazol-4-yl group, 2H-indazol-6-yl group, imidazo[1,2-a]pyridin-6-yl group, imidazo[1,2-a]pyridin-7-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1H-pyrrolo[2,3-b]pyridin-4-yl group, isoquinolin-6-yl group, isoquinolin-3-yl group, isoquinolin-5-yl group, isoquinolin-7-yl group, dihydro-2H-isoquinolin-6-yl group, dihydro-2H-isoquinolin-5-yl group, cinnolin-6-yl group, cinnolin-5-yl group, quinazolin-6-yl group, quinazolin-7-yl group, quinazolin-5-yl group, quinoxalin-2-yl group, quinoxalin-6-yl group, quinoxalin-5-yl group, 1H-benzimidazol-5-yl group, 1H-benzimidazol-4-yl group, benzoxazol-5-yl group, benzoxazol-6-yl group, benzoxazol-4-yl group, benzoxazol-7-yl group, 1H-pyrrolo[3,2-b]pyridin-5-yl group, 1H-pyrrolo[3,2-b]pyridin-6-yl group, benzo[1,2,5]thiadiazol-5-yl group, benzo[1,2,5]thiadiazol-4-yl group, 1H-benzotriazol-5-yl group, 1H-benzotriazol-4-yl group, 1,3-dihydropyrrolo[2,3-b]pyridin-5-yl group, 1,3-dihydropyrrolo[2,3-b]pyridin-4-yl group, 1,3-

dihydrobenzimidazol-5-yl group, 1,3-dihydrobenzimidazol-4-yl group, dihydro-3H-benzoxazol-6-yl group, dihydro-3H-benzoxazol-7-yl group, dihydro-3H-benzoxazol-5-yl group, dihydro-3H-benzoxazol-4-yl group, phthalazin-6-yl group, phthalazin-5-yl group, [1,8]naphthalidin-3-yl group, [1,8]naphthalidin-4-yl group, [1,5]naphthalidin-3-yl group, [1,5]naphthalidin-4-yl group, 1H-pyrrolo[3,2-c]pyridin-6-yl group, 1H-pyrrolo[3,2-c]pyridin-4-yl group, 1H-pyrrolo[2,3-c]pyridin-5-yl group, 1H-pyrrolo[2,3-c]pyridin-4-yl group, 1H-pyrazolo[4,3-b]pyridin-5-yl group, 1H-pyrazolo[4,3-b]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-4-yl group, 1H-pyrazolo[3,4-c]pyridin-5-yl group, 1H-pyrazolo[3,4-c]pyridin-4-yl group, 1H-pyrazolo[3,4-b]pyridin-5-yl group, 1H-pyrazolo[3,4-b]pyridin-4-yl group, [1,2,4]triazolo[4,3-a]pyridin-6-yl group, [1,2,4]triazolo[4,3-a]pyridin-7-yl group, thieno[3,2-c]pyridin-2-yl group, thieno[3,2-c]pyridin-3-yl group, thieno[3,2-c]pyridin-6-yl group, thieno[3,2-b]pyridin-2-yl group, thieno[3,2-b]pyridin-3-yl group, thieno[3,2-b]pyridin-5-yl group, thieno[3,2-b]pyridin-6-yl group, 1H-thieno[3,2-c]pyrazol-5-yl group, 1H-thieno[3,2-c]pyrazol-4-yl group, benzo[d]isoxazol-5-yl group, benzo[d]isoxazol-4-yl group, benzo[d]isoxazol-6-yl group, benzo[d]isoxazol-7-yl group, benzo[c]isoxazol-5-yl group, benzo[c]isoxazol-4-yl group, benzo[c]isoxazol-6-yl group, benzo[c]isoxazol-7-yl group, indolizin-7-yl group, indolizin-6-yl group, indolizine-8-yl group, 1,3-dihydroindol-5-yl group, 1,3-dihydroindol-4-yl group, 1,3-dihydroindol-6-yl group, 1H-pyrazolo[3,4-d]thiazol-5-yl group, 2H-isoindol-5-yl group, 2H-isoindol-4-yl group, [1,2,4]triazolo[1,5-a]pyrimidin-6-yl group, 1H-pyrazolo[3,4-b]pyrazin-5-yl group, 1H-imidazo[4,5-b]pyrazin-5-yl group, 7H-purin-2-yl group, 4H-chromen-6-yl group, or 4H-chromen-5-yl group (the aforementioned groups may be substituted with one of Xa or two or more of the same or different Xa), Xa is oxo group, thioxo group, fluorine atom, chlorine atom, trifluoromethyl group, methyl group, ethyl group, propyl group, 2-hydroxyethyl group, carboxymethyl group, 2-carboxyethyl group,

N,N-dimethylcarbamoylmethyl group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, carboxymethyloxy group, 2-carboxyethyloxy group, N,N-dimethylcarbamoylmethyloxy group, amino group, methylamino group, dimethylamino group, 2-hydroxyethylamino group, carbamoylamino group, acetylamino group, furan-2-carboxyamino group, 2-hydroxyacetylamino group, 2-aminoacetylamino group, methylsulfonylamino group, (N,N-dimethylsulfamoyl)amino group, methanesulfonyl group, sulfamoyl group, N-methylsulfamoyl group, N,N-dimethylsulfamoyl group, carboxyl group, acetyl group, carbamoyl group, or N,N-dimethylcarbamoyl group, and

Y is hydrogen atom, methyl group, or ethyl group.

(85) A medicament containing a compound represented by the aforementioned formula (I) or a pharmacologically acceptable salt thereof as an active ingredient.

(86) An agent for suppressing production of a prostaglandin and/or leukotriene, which contains a compound represented by the aforementioned formula (I) or a pharmacologically acceptable salt thereof as an active ingredient.

(87) The medicament according to (85) for prophylactic and/or therapeutic treatment of a disease caused by production of a prostaglandin and/or leukotriene.

(88) The medicament according to (85) for prophylactic and/or therapeutic treatment of an inflammatory disease of a mammal.

(89) The medicament according to (85) for prophylactic and/or therapeutic treatment of an autoimmune disease of a mammal.

(90) The medicament according to (85) for prophylactic and/or therapeutic treatment of an allergic disease of a mammal.

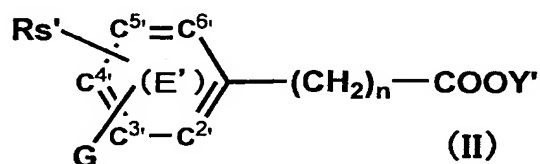
(91) The medicament according to (85) for defervescence and/or pain relief of a mammal.

(92) A pharmaceutical composition for prophylactic and/or therapeutic treatment of a condition of living body of a mammal exhibiting an acute or chronic inflammatory

reaction, which comprises a prophylactically and/or therapeutically effective amount of a compound represented by the aforementioned formula (I) or a pharmacologically acceptable salt thereof and a pharmaceutically acceptable carrier.

(93) A method for prophylactic and/or therapeutic treatment of a condition of living body of a mammal exhibiting an acute or chronic inflammatory reaction, which comprises administering a prophylactically and/or therapeutically effective amount of a compound represented by the aforementioned formula (I) or a pharmacologically acceptable salt thereof to the mammal.

(94) A compound represented by the following formula (II):



[In the formula, each of C<sup>2'</sup>, C<sup>3'</sup>, C<sup>4'</sup>, C<sup>5'</sup>, and C<sup>6'</sup> in the aromatic ring (E') represents a ring-constituting carbon atom, any one of them to which Rs' and G does not bind may be replaced with V',

V' represents nitrogen atom, or carbon atom substituted with Zx', Zx' has the same meaning as Zx mentioned above, provided that when Zx contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when Zx contains amino group, the amino group may be protected with Rp<sup>2</sup>,

Rs' represents -D-Rx' or -N(Ry')(Rz'),

-D-Rx' and -N(Ry')(Rz') have the same meanings as -D-Rx and -N(Ry)(Rz), respectively, provided that when -D-Rx or -N(Ry)(Rz) contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, when -D-Rx or -N(Ry)(Rz) contains amino group, the amino group may be protected with Rp<sup>2</sup>,

G represents chlorine atom, bromine atom, iodine atom, mesylate group,

triflate group, or an arenesulfonate group of which aromatic portion may be substituted with one of T<sup>1</sup> or two or more of the same or different T<sup>1</sup>, and

Y' represents a lower alkyl group having 1 to 4 carbon atoms].

(95) The compound according to (94) mentioned above, wherein, in the formula (II), G binds to the ring-constituting carbon atom C<sup>2'</sup> or C<sup>3'</sup> in the aromatic ring (E').

(96) The compound according to (94) or (95) mentioned above, wherein, in the formula (II), n is an integer of 2.

(97) The compound according to any one of (94) to (96) mentioned above, wherein, in the formula (II), Rs' is -O-Rx'.

(98) The compound according to any one of (94) to (97) mentioned above, wherein, in the formula (II), Rs' is -D-Rx' or -N(Ry')(Rz'), D is a single bond, oxygen atom, sulfur atom, -S(O)-, -S(O)<sub>2</sub>-, or -C(O)-, Rx' is a linear or branched saturated alkyl group having 3 to 8 carbon atoms, or Ra, Rb, or Rc, k in Ra is 0 or an integer of 1 to 3, R<sup>1</sup> is a saturated cyclic alkyl group having 3 to 7 carbon atoms or a condensed saturated cyclic alkyl group having 6 to 8 carbon atoms, R<sup>1</sup> may be substituted with one of lower alkyl group having 1 to 4 carbon atoms or two or more of the same or different lower alkyl groups having 1 to 4 carbon atoms, Q in Rb is phenyl group, thienyl group, furyl group, pyrrolyl group, pyridyl group, oxazolyl group, isoxazolyl group, thiazolyl group, isothiazolyl group, imidazolyl group, pyrazolyl group, oxadiazolyl group, thiadiazolyl group, triazolyl group, tetrazolyl group, naphthyl group, tetrahydronaphthyl group, indanyl group, indenyl group, quinolyl group, isoquinolyl group, indolyl group, benzofuryl group, benzothienyl group, benzimidazolyl group, benzoxazolyl group, benzothiazolyl group, indazolyl group, 4H-chromenyl group, dihydrobenzodioxyl group, benzoisoxazolyl group, pyrrolopyridinyl group, pyrazolopyridinyl group, triazolopyridinyl group, thienopyridinyl group, thienopyrazolyl group, 1,3-dihydrobenzimidazole group, dihydro-3H-benzoxazole group, or dihydro-3H-benzothiazole group, which binds to

$A^2$  at an arbitrary position on the ring,  $A^1$  is a single bond or an alkylene (a) having 1 to 3 carbon atoms, the alkylene (a) may be substituted with a lower alkyl group having 1 to 4 carbon atoms or phenyl group,  $A^2$  is a single bond, oxygen atom, sulfur atom,  $-S(O)-$ ,  $-S(O)_2-$ , or  $-N(R^4)-$  (provided that when  $A^2$  is oxygen atom, sulfur atom,  $-S(O)-$ ,  $-S(O)_2-$ , or  $-N(R^4)-$ ,  $A^1$  is ethylene or trimethylene),  $R^2$  and  $R^3$  independently represent hydrogen atom, a linear or branched saturated alkyl group having 1 to 4 carbon atoms, oxo group, thioxo group, fluorine atom, chlorine atom, bromine atom, trifluoromethyl group,  $-OR^5$ ,  $-N(R^6)(R^6)$ ,  $-NHCOR^7$ ,  $-NHSO_2R^8$ , or  $-A^6-Qa$ , or they bind to each other to form methylenedioxy group,  $Qa$  may be substituted with one of  $T^1$  or two or more of the same or different  $T^1$ , and is phenyl group, pyridyl group, oxazolyl group, isoxazolyl group, thiazolyl group, isothiazolyl group, imidazolyl group, pyrazolyl group, oxadiazolyl group, thiadiazolyl group, triazolyl group, tetrazolyl group, naphthyl group, indanyl group, indenyl group, quinolyl group, isoquinolyl group, indolyl group, benzofuryl group, benzothienyl group, benzimidazolyl group, benzoxazolyl group, benzothiazolyl group, or indazolyl group, which binds to  $A^6$  at an arbitrary position on the ring,  $R^4$  and  $R^6$  independently represent hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms,  $R^5$  and  $R^7$  independently represent hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms, or  $-A^6-Qa$ ,  $R^8$  represents a lower alkyl group having 1 to 4 carbon atoms,  $R^6$  has the same meaning as  $R^6$ , or binds to  $R^6$  to form a 3- to 6-membered ring of a cycloalkyl group or morpholino group together with the nitrogen atom to which they bind,  $p$  in  $R_c$  is an integer of 2 to 4,  $A^4$  is a single bond or methylene or ethylene,  $A^5$  is  $-C(O)-$ ,  $-C(S)-$ , or  $-S(O)_2-$ ,  $R_d$  is hydrogen atom, an alkyl group having 1 to 8 carbon atoms, or  $Qa$ ,  $R_e$  is an alkyl group having 1 to 8 carbon atoms,  $-A^6-Qa$ ,  $-(CH_2)_iR^{14}$ ,  $-OR^{28}$ ,  $-SR^{28}$ , or  $-N(R^{29})(R^{30})$ ,  $i$  is an integer of 1 to 3,  $R^{14}$  is hydroxyl group, an alkoxy group having 1 to 4 carbon atoms, carboxyl group, or an  $N,N$ -dialkylcarbamoyl group having 1 to 4 carbon atoms,  $R^{28}$  is an alkyl group having 1

to 8 carbon atoms or  $-A^6-Qa$ ,  $R^{29}$  is an alkyl group having 1 to 8 carbon atoms, an alkoxy carbonyl group having 1 to 4 carbon atoms, or  $-A^6-Qa$ ,  $R^{30}$  represents hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms, or binds to  $R^{29}$  to form a 3- to 6-membered ring of nitrogen-containing cycloalkyl group or morpholino group together with the nitrogen atom to which they bind,  $Rz'$  has the same meaning as  $Rx'$ , or represents  $-A^5-Re$ ,  $Ry'$  represents hydrogen atom, an alkyl group having 1 to 8 carbon atoms, or  $-A^6-Qp$ , or binds to  $Rz'$  to form a saturated or unsaturated nitrogen-containing cyclic substituent having 3 to 7 atoms together with the nitrogen atom to which they bind, when  $-D-Rx'$  or  $-N(Ry')(Rz')$  contains hydroxyl group, the hydroxyl group may be protected with  $Rp^1$ , and when  $-D-Rx'$  or  $-N(Ry')(Rz')$  contains amino group, the amino group may be protected with  $Rp^2$ .

(99) The compound according to (94) mentioned above, wherein, in the formula (II),  $n$  is an integer of 1 to 3,

$G$  binds to  $C^{3'}$ ,  $Rs'$  binds to one of the ring-constituting carbon atoms  $C^{4'}$ ,  $C^{5'}$ , and  $C^{6'}$ , a ring-constituting carbon atom to which  $Rs'$  does not bind among  $C^{4'}$ ,  $C^{5'}$ , and  $C^{6'}$  may be replaced with  $V'$ ,

$V'$  is nitrogen atom or carbon atom substituted with  $Zx'$ ,  $Zx'$  is fluorine atom, chlorine atom, bromine atom, nitro group, methyl group, hydroxyl group, methoxy group, amino group, N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, or N,N-dimethylsulfamoylamino group, provided that when  $Zx'$  contains hydroxyl group, the hydroxyl group may be protected with  $Rp^1$ , and when  $Zx'$  contains amino group, the amino group may be protected with  $Rp^2$ ,

$Rs'$  is  $-D-Rx'$  or  $-N(Ry')(Rz')$ ,  $D$  is oxygen atom or sulfur atom,  $Rx'$  is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-



cyclopentylethyl group, or 2-cyclohexylethyl group, or Rb or Rc, Q in Rb is phenyl group, thienyl group, furyl group, pyridyl group, oxazolyl group, naphthyl group, tetrahydronaphthyl group, indanyl group, indolyl group, or dihydrobenzodioxyl group, A<sup>2</sup> is a single bind, oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)- (provided that when A<sup>2</sup> is oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)-, A<sup>1</sup> is ethylene), R<sup>2</sup> and R<sup>3</sup> independently represent hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, dimethylamino group, acetylamino group, or methylsulfonylamino group (provided that when Q is phenyl group, A<sup>1</sup> is a single bind or unsubstituted methylene, and A<sup>2</sup> is a single bind, one of R<sup>2</sup> and R<sup>3</sup> is a substituent other than hydrogen atom), p in Rc is an integer of 2 or 3, A<sup>4</sup> is a single bind or methylene, A<sup>5</sup> is -C(O)-, -C(S)-, or -S(O)<sub>2</sub>-, Rd is hydrogen atom, or methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group, cyclopropylmethyl group, cyclopentyl group, cyclopentylmethyl group, cyclohexyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, or pyridin-4-yl group, Re is methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, phenylmethyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, furan-2-yl group, furan-3-yl group, thiophen-2-yl group, thiophen-3-yl group, methoxy group, ethoxy group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenyloxy group, 4-

methylphenoxy group, 4-chlorophenoxy group, 4-fluorophenoxy group, thiomethoxy group, amino group, N-methylamino group, N,N-dimethylamino group, N-ethylamino group, N,N-diethylamino group, N-propylamino group, N-isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group, N-(4-fluorophenyl)amino group, N-(pyridin-2-yl)amino group, N-(pyridin-3-yl)amino group, N-(pyridin-4-yl)amino group, N-(furan-2-yl)amino group, N-(furan-3-yl)amino group, N-(thiophen-2-yl)amino group, N-(thiophen-3-yl)amino group, pyrrolidino group, piperidino group, morpholino group, methoxycarbonylamino group or ethoxycarbonylamino group, R<sub>z</sub> is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-

chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, 2-(N-ethyl-N-phenylamino)ethyl group, isobutyryl group, isopropylthiocarbonyl group, isopropylsulfonyl group, valeryl group, butylthiocarbonyl group, isovaleryl group, isobutylthiocarbonyl group, pivaloyl group, t-butylthiocarbonyl group, cyclopropylcarbonyl group, cyclopropylthiocarbonyl group, cyclopentylcarbonyl group, cyclopentylthiocarbonyl group, cyclohexylcarbonyl group, cyclohexylthiocarbonyl group, cyclopentylmethylcarbonyl group, cyclopentylmethylthiocarbonyl group, cyclohexylmethylcarbonyl group, cyclohexylmethylthiocarbonyl group, benzoyl group, thiobenzoyl group, phenylsulfonyl group, 4-methylphenylcarbonyl group, 4-methylphenylthiocarbonyl group, 4-methylphenylsulfonyl group, 4-

chlorophenylcarbonyl group, 4-chlorophenylthiocarbonyl group, 4-fluorophenylcarbonyl group, 4-fluorophenylthiocarbonyl group, isopropylloxycarbonyl group, N-isopropylcarbamoyl group, N-isopropylthiocarbamoyl group, butylloxycarbonyl group, N-butylcarbamoyl group, N-butylthiocarbamoyl group, isobutylloxycarbonyl group, N-isobutylcarbamoyl group, N-isobutylthiocarbamoyl group, t-butylloxycarbonyl group, N-t-butylcarbamoyl group, N-t-butylthiocarbamoyl group, cyclopropylloxycarbonyl group, N-cyclopropylcarbamoyl group, N-cyclopropylthiocarbamoyl group, cyclopentylloxycarbonyl group, N-cyclopentylcarbamoyl group, N-cyclopentylthiocarbamoyl group, cyclohexylloxycarbonyl group, N-cyclohexylcarbamoyl group, N-cyclohexylthiocarbamoyl group, cyclopentylmethyloxycarbonyl group, cyclohexylmethyloxycarbonyl group, phenyloxycarbonyl group, N-phenylcarbamoyl group, N-phenylthiocarbamoyl group, 4-methylphenyloxycarbonyl group, N-(4-methylphenyl)carbamoyl group, N-(4-methylphenyl)thiocarbamoyl group, 4-chlorophenyloxycarbonyl group, N-(4-chlorophenyl)carbamoyl group, N-(4-chlorophenyl)thiocarbamoyl group, 4-fluorophenyloxycarbonyl group, N-(4-fluorophenyl)carbamoyl group, N-(4-fluorophenyl)thiocarbamoyl group, (pyrrolidino-1-yl)carbonyl group, (piperidino-1-yl)carbonyl group, or (morpholino-4-yl)carbonyl group, Ry is hydrogen atom, methyl group, ethyl group or isobutyl group, or binds to Rz to form pyrrolidino group, piperidino group, piperazino group, morpholino group, pyrrol-1-yl group, imidazol-1-yl group, or pyrazol-1-yl group together with the nitrogen atom to which they binds, provided that when -D-Rx' or -N(Ry')(Rz') contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when -D-Rx' or -N(Ry')(Rz') contains amino group, the amino group may be protected with Rp<sup>2</sup>,

G is chlorine atom, bromine atom, iodine atom, or triflate group, and

Y' is methyl group or ethyl group.

(100) The compound according to any one of (94) to (96) mentioned above, wherein, in the formula (II),  $R_{s'}$  is  $-N(R_{y'})(R_{z'})$ .

(101) The compound according to any one of (94) to (96) mentioned above, wherein, in the formula (II),  $R_{s'}$  is  $-D-R_{x'}$ , and D is sulfur atom,  $-S(O)-$ ,  $-S(O)_2-$  or  $-C(O)-$ .

(102) The compound according to (94) mentioned above, wherein, in the formula (II), G binds at the position of  $C^{2'}$  in the aromatic ring ( $E'$ ),  $R_{s'}$  binds to one of the ring-constituting carbon atoms  $C^{3'}$ ,  $C^{4'}$  and  $C^{5'}$ , and all of  $C^{2'}$ ,  $C^{3'}$ ,  $C^{4'}$ ,  $C^{5'}$  and  $C^{6'}$  in the aromatic ring ( $E'$ ) are not replaced with  $V'$ .

(103) The compound according to (94) mentioned above, wherein, in the formula (II), n is an integer of 1 to 3,

G binds to  $C^{2'}$ ,  $R_{s'}$  binds to one of the ring-constituting carbon atoms  $C^{4'}$ ,  $C^{5'}$ , and  $C^{6'}$ , a ring-constituting carbon atom to which  $R_{s'}$  does not bind among  $C^{3'}$ ,  $C^{4'}$ , and  $C^{5'}$  may be replaced with  $V'$ ,

$V'$  is nitrogen atom, or carbon atom substituted with  $Zx'$ ,  $Zx'$  is fluorine atom, chlorine atom, bromine atom, nitro group, methyl group, hydroxyl group, methoxy group, amino group, N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, or N,N-dimethylsulfamoylamino group, provided that when  $Zx'$  contains hydroxyl group, the hydroxyl group may be protected with  $R_{p^1}$ , and when  $Zx'$  contains amino group, the amino group may be protected with  $R_{p^2}$ ,

$R_{s'}$  is  $-D-R_{x'}$  or  $-N(R_{y'})(R_{z'})$ , D is oxygen atom or sulfur atom,  $R_{x'}$  is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-cyclopentylethyl group, 2-cyclohexylethyl group, or Rb or Rc, Q in Rb is phenyl group, thienyl group, furyl group, pyridyl group, oxazolyl group, naphthyl group, tetrahydronaphthyl group, indanyl group, indolyl group, or dihydrobenzodioxyl

group, A<sup>2</sup> is a single bind, oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)- (provided that when A<sup>2</sup> is oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)-, A<sup>1</sup> is ethylene), R<sup>2</sup> and R<sup>3</sup> independently represent hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, dimethylamino group, acetylamino group, or methylsulfonylamino group (provided that when Q is phenyl group, A<sup>1</sup> is a single bind or unsubstituted methylene, and A<sup>2</sup> is a single bind, one of R<sup>2</sup> and R<sup>3</sup> is a substituent other than hydrogen atom), p in R<sub>c</sub> is an integer of 2 or 3, A<sup>4</sup> is a single bind or methylene, A<sup>5</sup> is -C(O)-, -C(S)-, or -S(O)<sub>2</sub>-, R<sub>d</sub> is hydrogen atom, or methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group, cyclopropylmethyl group, cyclopentyl group, cyclopentylmethyl group, cyclohexyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, or pyridin-4-yl group, R<sub>e</sub> is methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, phenylmethyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, furan-2-yl group, furan-3-yl group, thiophen-2-yl group, thiophen-3-yl group, methoxy group, ethoxy group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenyloxy group, 4-methylphenyloxy group, 4-chlorophenyloxy group, 4-fluorophenyloxy group, thiomethoxy group, amino group, N-methylamino group, N,N-dimethylamino group, N-ethylamino group, N,N-diethylamino group, N-propylamino group, N-

isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group, N-(4-fluorophenyl)amino group, N-(pyridin-2-yl)amino group, N-(pyridin-3-yl)amino group, N-(pyridin-4-yl)amino group, N-(furan-2-yl)amino group, N-(furan-3-yl)amino group, N-(thiophen-2-yl)amino group, N-(thiophen-3-yl)amino group, pyrrolidino group, piperidino group, morpholino group, methyloxycarbonylamino group, or ethyloxycarbonylamino group, R<sub>z</sub> is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-

dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, 2-(N-ethyl-N-phenylamino)ethyl group, isobutyryl group, isopropylthiocarbonyl group, isopropylsulfonyl group, valeryl group, butylthiocarbonyl group, isovaleryl group, isobutylthiocarbonyl group, pivaloyl group, t-butylthiocarbonyl group, cyclopropylcarbonyl group, cyclopropylthiocarbonyl group, cyclopentylcarbonyl group, cyclopentylthiocarbonyl group, cyclohexylcarbonyl group, cyclohexylthiocarbonyl group, cyclopentylmethylcarbonyl group, cyclopentylmethylthiocarbonyl group, cyclohexylmethylcarbonyl group, cyclohexylmethylthiocarbonyl group, benzoyl group, thiobenzoyl group, phenylsulfonyl group, 4-methylphenylcarbonyl group, 4-methylphenylthiocarbonyl group, 4-methylphenylsulfonyl group, 4-chlorophenylcarbonyl group, 4-chlorophenylthiocarbonyl group, 4-fluorophenylcarbonyl group, 4-fluorophenylthiocarbonyl group, isopropoxy carbonyl group, N-isopropylcarbamoyl group, N-isopropylthiocarbamoyl



group, butyloxycarbonyl group, N-butylcarbamoyl group, N-butylthiocarbamoyl group, isobutyloxycarbonyl group, N-isobutylcarbamoyl group, N-isobutylthiocarbamoyl group, t-butyloxycarbonyl group, N-t-butylcarbamoyl group, N-t-butylthiocarbamoyl group, cyclopropyloxycarbonyl group, N-cyclopropylcarbamoyl group, N-cyclopropylthiocarbamoyl group, cyclopentyloxycarbonyl group, N-cyclopentylcarbamoyl group, N-cyclopentylthiocarbamoyl group, cyclohexyloxycarbonyl group, N-cyclohexylcarbamoyl group, N-cyclohexylthiocarbamoyl group, cyclopentylmethyloxycarbonyl group, cyclohexylmethyloxycarbonyl group, phenyloxycarbonyl group, N-phenylcarbamoyl group, N-phenylthiocarbamoyl group, 4-methylphenyloxycarbonyl group, N-(4-methylphenyl)carbamoyl group, N-(4-methylphenyl)thiocarbamoyl group, 4-chlorophenyloxycarbonyl group, N-(4-chlorophenyl)carbamoyl group, N-(4-chlorophenyl)thiocarbamoyl group, 4-fluorophenyloxycarbonyl group, N-(4-fluorophenyl)carbamoyl group, N-(4-fluorophenyl)thiocarbamoyl group, (pyrrolidino-1-yl)carbonyl group, (piperidino-1-yl)carbonyl group, or (morpholino-4-yl)carbonyl group,  $R_y$  is hydrogen atom, methyl group, ethyl group or isobutyl group, or binds to  $R_z$  to form pyrrolidino group, piperidino group, piperazino group, morpholino group, pyrrol-1-yl group, imidazol-1-yl group, or pyrazol-1-yl group together with the nitrogen atom, provided that when  $-D-R_x'$  or  $-N(R_y')(R_z')$  contains hydroxyl group, the hydroxyl group may be protected with  $R_p^1$ , and when  $-D-R_x'$  or  $-N(R_y')(R_z')$  contains amino group, the amino group may be protected with  $R_p^2$ ,

$G$  is chlorine atom, bromine atom, iodine atom, or triflate group, and

$Y'$  is methyl group or ethyl group.

(104) The compound according to (102) or (103) mentioned above, wherein, in the formula (II),  $n$  is an integer of 2,  $Rs'$  binds to  $C^{3'}$  in the aromatic ring ( $E'$ ),  $Rs'$  is  $-O-R_x'$ , and  $Y'$  is methyl group or ethyl group.

(105) The compound according to (94) mentioned above, wherein, in the formula (II),  $n$  is an integer of 2,

$C^2$  is a ring-constituting carbon atom to which  $G$  binds,  $C^3$  is a ring-constituting carbon atom to which  $Rs'$  binds,  $C^4$  may be replaced with  $V$ ,  $C^5$  and  $C^6$  are unsubstituted ring-constituting carbon atoms,

$V$  is nitrogen atom, or carbon atom substituted with  $Zx'$ ,  $Zx'$  is fluorine atom, methyl group, hydroxyl group, amino group,  $N$ -methylamino group, or  $N,N$ -dimethylamino group, provided that when  $Zx'$  contains hydroxyl group, the hydroxyl group may be protected with  $Rp^1$ , and when  $Zx'$  contains amino group, the amino group may be protected with  $Rp^2$ ,

$Rs'$  is  $-O-Rx'$ ,  $Rx'$  is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-

chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, or 2-(N-ethyl-N-phenylamino)ethyl group,

G is bromine atom or iodine atom, and

Y' is methyl group or ethyl group.

(106) The compound according to (102) or (103) mentioned above, wherein, in the formula (II), n is an integer of 2, Rs' binds at the position of C<sup>4'</sup> in the aromatic ring (E'), Rs' is -O-Rx', and Y' is methyl group or ethyl group.

(107) The compound according to (94) mentioned above, wherein, in the formula (II), n is an integer of 2,

C<sup>2'</sup> is a ring-constituting carbon atom to which G binds, C<sup>4'</sup> is a ring-

constituting carbon atom to which Rs' binds, C<sup>5'</sup> may be replaced with V', C<sup>3'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms,

V' is nitrogen atom, or carbon atom substituted with Zx', Zx' is fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, or N,N-dimethylamino group, provided that when Zx' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when Zx' contains amino group, the amino group may be protected with Rp<sup>2</sup>,

Rs' is -O-Rx', Rx' is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-

dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, or 2-(N-ethyl-N-phenylamino)ethyl group,

G is bromine atom or iodine atom, and

Y' is methyl group or ethyl group.

(108) The compound according to (102) or (103) mentioned above, wherein, in the formula (II), n is an integer of 2, Rs' binds at the position of C<sup>5'</sup> in the aromatic ring (E'), Rs' is -O-Rx', and Y' is methyl group or ethyl group.

(109) The compound according to (94) mentioned above, wherein, in the formula (II), G binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>5'</sup> or C<sup>6'</sup> in the aromatic ring (E'), and all of C<sup>2'</sup>, C<sup>3'</sup>, C<sup>4'</sup>, C<sup>5'</sup> and C<sup>6'</sup> in the aromatic ring (E') are not replaced with V'.

(110) The compound according to (109) mentioned above, wherein, in the formula (II), n is an integer of 2, Rs' binds to C<sup>5'</sup> in the aromatic ring (E'), and Rs' is -O-Rx'.

(111) The compound according to (94) mentioned above, wherein, in the formula (II),  $n$  is an integer of 2,

$C^{3'}$  is carbon atom to which  $G$  binds,  $C^{5'}$  is carbon atom to which  $Rs'$  binds,  $C^{2'}$ ,  $C^{4'}$ , and  $C^{6'}$  are unsubstituted ring-constituting carbon atoms,

$Rs'$  is  $-O-Rx'$ ,  $Rx'$  is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-

(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group or 2-(N-ethyl-N-phenylamino)ethyl group,

G is bromine atom or iodine atom, and

Y' is methyl group or ethyl group.

(112) The compound according to (109) mentioned above, wherein, in the formula

(II), n is an integer of 2, Rs' binds to C<sup>6'</sup> in the aromatic ring (E'), and Rs' is -O-Rx'.

(113) The compound according to (94) mentioned above, wherein, in the formula (II),

G binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), and C<sup>6'</sup> is V'.

(114) The compound according to (113) mentioned above, wherein, in the formula

(II), n is an integer of 2, V' is carbon atom substituted with Zx', and Rs' is -O-Rx'.

(115) The compound according to (94) mentioned above, wherein, in the formula (II),

n is an integer of 2,

C<sup>3'</sup> is carbon atom to which G binds, C<sup>4'</sup> is a carbon atom to which Rs' binds, C<sup>6'</sup> is carbon atom substituted with Zx', C<sup>2'</sup> and C<sup>5'</sup> are unsubstituted ring-constituting carbon atoms,

Zx' is fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, or N,N-dimethylamino group, provided that when Zx' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when Zx' contains amino group, the amino group may be protected with Rp<sup>2</sup>,

Rs' is -O-Rx', Rx' is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-



(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, or 2-(N-ethyl-N-phenylamino)ethyl group,

G is bromine atom or iodine atom, and

Y' is methyl group or ethyl group.

(116) The compound according to (94) mentioned above, wherein, in the formula (II), G binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5'</sup> is nitrogen atom, and C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms.

(117) The compound according to (116) mentioned above, wherein, in the formula (II), n is an integer of 2, and Rs' is -O-Rx'.

(118) The compound according to (94) mentioned above, wherein, in the formula (II), n is an integer of 2,

C<sup>3'</sup> is carbon atom to which G binds, C<sup>4'</sup> is carbon atom to which Rs' binds, C<sup>5'</sup> is nitrogen atom, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms,

Rs' is -O-Rx', Rx' is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-

fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-

(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, or 2-(N-ethyl-N-phenylamino)ethyl group,

G is bromine atom or iodine atom, and

Y' is methyl group or ethyl group.

(119) The compound according to (94) mentioned above, wherein, in the formula (II), G binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5'</sup> is a ring-constituting carbon atom substituted with Zx', or an unsubstituted ring-constituting carbon atom, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms, Rs' is -D-Rx', and D is a single bond, sulfur atom, -S(O)-, -S(O)<sub>2</sub>-, or -C(O)-.

(120) The compound according to (94) mentioned above, wherein, in the formula (II), n is an integer of 2,

C<sup>3'</sup> is carbon atom to which G binds, C<sup>4'</sup> is a carbon atom to which Rs' binds, C<sup>5'</sup> is a ring-constituting carbon atom substituted with Zx', or an unsubstituted ring-constituting carbon atom, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms,

Zx' is fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, or N,N-dimethylamino group, provided that when Zx' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when Zx' contains amino group, the amino group may be protected with Rp<sup>2</sup>,

Rs' is -S-Rx', Rx' is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl

group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-

(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, or 2-(N-ethyl-N-phenylamino)ethyl group,

G is bromine atom or iodine atom, and

Y' is methyl group or ethyl group.

(121) The compound according to (94) mentioned above, wherein, in the formula (II), G binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5'</sup> is a ring-constituting carbon atom substituted with Zx', or an unsubstituted ring-constituting carbon atom, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms, and Rs' is -N(Ry')(Rz').

(122) The compound according to (94) mentioned above, wherein, in the formula (II), n is an integer of 2,

C<sup>3'</sup> is carbon atom to which G binds, C<sup>4'</sup> is a carbon atom to which Rs' binds, C<sup>5'</sup> is a ring-constituting carbon atom substituted with Zx', or an unsubstituted ring-constituting carbon atom, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms,

Zx' is fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, or N,N-dimethylamino group, provided that when Zx' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when Zx' contains amino group, the amino group may be protected with Rp<sup>2</sup>,

Rs' is -N(Ry')(Rz'), Rz' is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-

methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group,  
 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group,  
 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group,  
 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group,  
 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl  
 group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl  
 group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-  
 chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl  
 group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-  
 methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-  
 dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl  
 group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-  
 chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl  
 group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-  
 difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-  
 dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-  
 dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-  
 dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-  
 (trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-  
 (trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-  
 methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-  
 methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-  
 methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl  
 group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-  
 chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-  
 (trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-  
 (trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group,

2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, 2-(N-ethyl-N-phenylamino)ethyl group, isobutyryl group, isopropylthiocarbonyl group, isopropylsulfonyl group, valeryl group, butylthiocarbonyl group, isovaleryl group, isobutylthiocarbonyl group, pivaloyl group, t-butylthiocarbonyl group, cyclopropylcarbonyl group, cyclopropylthiocarbonyl group, cyclopentylcarbonyl group, cyclopentylthiocarbonyl group, cyclohexylcarbonyl group, cyclohexylthiocarbonyl group, cyclopentylmethylcarbonyl group, cyclopentylmethylthiocarbonyl group, cyclohexylmethylcarbonyl group, cyclohexylmethylthiocarbonyl group, benzoyl group, thiobenzoyl group, phenylsulfonyl group, 4-methylphenylcarbonyl group, 4-methylphenylthiocarbonyl group, 4-methylphenylsulfonyl group, 4-chlorophenylcarbonyl group, 4-chlorophenylthiocarbonyl group, 4-fluorophenylcarbonyl group, 4-fluorophenylthiocarbonyl group, isopropoxy carbonyl group, N-isopropylcarbamoyl group, N-isopropylthiocarbamoyl group, butyloxy carbonyl group, N-butylcarbamoyl group, N-butylthiocarbamoyl group, isobutyloxy carbonyl group, N-isobutylcarbamoyl group, N-isobutylthiocarbamoyl group, t-butyloxy carbonyl group, N-t-butylcarbamoyl group, N-t-butylthiocarbamoyl group, cyclopropyloxy carbonyl group, N-cyclopropylcarbamoyl group, N-cyclopropylthiocarbamoyl group, cyclopentyloxy carbonyl group, N-cyclopentylcarbamoyl group, N-cyclopentylthiocarbamoyl group, cyclohexyloxy carbonyl group, N-cyclohexylcarbamoyl group, N-cyclohexylthiocarbamoyl group, cyclopentylmethyloxy carbonyl group, cyclohexylmethyloxy carbonyl group, phenyloxy carbonyl group, N-phenylcarbamoyl group, N-phenylthiocarbamoyl group, 4-methylphenyloxy carbonyl group, N-(4-methylphenyl)carbamoyl group, N-(4-methylphenyl)thiocarbamoyl group, 4-chlorophenyloxy carbonyl group, N-(4-

chlorophenyl)carbamoyl group, N-(4-chlorophenyl)thiocarbamoyl group, 4-fluorophenyloxycarbonyl group, N-(4-fluorophenyl)carbamoyl group, N-(4-fluorophenyl)thiocarbamoyl group, (pyrrolidino-1-yl)carbonyl group, (piperidino-1-yl)carbonyl group, or (morpholino-4-yl)carbonyl group, Ry' is hydrogen atom, methyl group, ethyl group or isobutyl group, or binds to Rz' to form pyrrolidino group, piperidino group, or morpholino group together with nitrogen atom to which they binds, provided that when -N(Ry')(Rz') contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when -N(Ry')(Rz') contains amino group, the amino group may be protected with Rp<sup>2</sup>,

G is bromine atom or iodine atom, and

Y' is methyl group or ethyl group.

(123) The compound according to (94) mentioned above, wherein, in the formula (II), n is an integer of 2, G binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5'</sup> is carbon atom substituted with -N(Rn<sup>1</sup>)(Rn<sup>2</sup>) group (provided that one of Rn<sup>1</sup> and Rn<sup>2</sup> is a substituent other than hydrogen atom), C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms, and Rs' is -O-Rx'.

(124) The compound according to (94) mentioned above, wherein, in the formula (II), n is an integer of 2,

C<sup>3'</sup> is carbon atom to which G binds, C<sup>4'</sup> is carbon atom to which Rs' binds, C<sup>5'</sup> is carbon atom substituted with Zx', C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms,

Zx' is N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, or N,N-dimethylsulfamoylamino group, provided that when Zx' contains amino group, the amino group may be protected with Rp<sup>2</sup>,

Rs' is -O-Rx', Rx' is butyl group, isobutyl group, 2-ethylbutyl group,



cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group,  
 cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-  
 fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl  
 group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-  
 methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group,  
 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group,  
 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group,  
 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group,  
 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl  
 group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl  
 group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-  
 chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl  
 group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-  
 methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-  
 dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl  
 group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-  
 chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl  
 group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-  
 difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-  
 dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-  
 dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-  
 dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-  
 (trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-  
 (trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-  
 methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-  
 methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-  
 methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl

group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, or 2-(N-ethyl-N-phenylamino)ethyl group,

G is bromine atom or iodine atom, and

Y' is methyl group or ethyl group.

(125) The compound according to (94) mentioned above, wherein, in the formula (II), G binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5'</sup> is a ring-constituting carbon atom substituted with Zx', or an unsubstituted ring-constituting carbon atom, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms, Rs' is -D-Rx', and Rx' has the same meaning as Rc, provided that when Rc contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when Rc contains amino group, the amino group may be protected with Rp<sup>2</sup>.

(126) The compound according to (94) mentioned above, wherein, in the formula (II), n is an integer of 2,

C<sup>3'</sup> is carbon atom to which G binds, C<sup>4'</sup> is a carbon atom to which Rs' binds, C<sup>5'</sup> is a ring-constituting carbon atom substituted with Zx', or an unsubstituted ring-constituting carbon atom, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms,

Zx' is fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, or N,N-dimethylamino group, provided that when Zx' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when Zx' contains amino group, the amino group may be protected with Rp<sup>2</sup>,

Rs' is -O-Rx', Rx' has the same meaning as Rc, provided that when Rc contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, p in Rc is an integer of 2, A<sup>4</sup> is a single bond or methylene, A<sup>5</sup> is -C(O)-, -C(S)-, or -S(O)<sub>2</sub>-, Rd is methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, or 4-fluorophenylmethyl group, Re is isopropyl group, butyl group, isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenyloxy group, 4-methylphenyloxy group, 4-chlorophenyloxy group, 4-fluorophenyloxy group, N-propylamino group, N-isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group, N-(4-fluorophenyl)amino group, pyrrolidino group, piperidino group, or morpholino group,

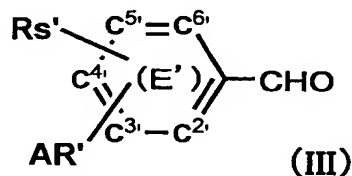
G is bromine atom or iodine atom, and

Y' is methyl group or ethyl group.

(127) The compound according to (94) mentioned above, wherein, in the formula (II), G binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5'</sup> is a ring-constituting carbon atom substituted with Zx', or an unsubstituted ring-constituting carbon atom, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms, Rs' is -O-Rx', and Rx' is a linear or branched saturated alkyl group having 3 to 8 carbon atoms, or Ra or Rb.

(128) The compound according to (94) mentioned above, wherein, in the formula (II),  $n$  is an integer of 2,  $G$  binds to  $C^{3'}$  in the aromatic ring ( $E'$ ),  $Rs'$  binds to  $C^{4'}$  in the aromatic ring ( $E'$ ),  $C^{5'}$  is carbon atom substituted with nitro group,  $C^{2'}$  and  $C^{6'}$  are unsubstituted ring-constituting carbon atoms, and  $Rs'$  is  $-O-Rx'$ .

(129) A compound represented by the following formula (III):



[In the formula,  $C^{2'}$ ,  $C^{3'}$ ,  $C^{4'}$ ,  $C^{5'}$  and  $C^{6'}$  in the aromatic ring ( $E'$ ) represent a ring-constituting carbon atom, any one of these atoms to which  $Rs'$  and  $AR'$  does not bind may be replaced with  $V'$ , and  $AR'$  has the same meaning as that of  $AR$ , provided that when  $AR$  contains hydroxyl group, the hydroxyl group may be protected with  $Rp^1$ , and when  $AR$  contains amino group, the amino group may be protected with  $Rp^2$ .].

(130) The compound according to (129) mentioned above, wherein, in the formula (III),  $AR'$  binds to the atom of  $C^{2'}$  or  $C^{3'}$  in the aromatic ring ( $E'$ ).

(131) The compound according to (129) or (130) mentioned above, wherein, in the formula (III),  $AR'$  is a residue of naphthalene, benzofuran, benzo[b]thiophene, indole, benzothiazole, dihydro-3H-benzothiazole, quinoline, dihydro-1H-quinoline, benzo[d]isothiazole, 1H-indazole, benzo[c]isothiazole, 2H-indazole, imidazo[1,2-a]pyridine, 1H-pyrrolo[2,3-b]pyridine, isoquinoline, dihydro-2H-isoquinoline, cinnoline, quinazoline, quinoxaline, 1H-benzimidazole, benzoxazole, 1H-pyrrolo[3,2-b]pyridine, benzo[1,2,5]thiadiazole, 1H-benzotriazole, 1,3-dihydropyrrolo[2,3-b]pyridine, 1,3-dihydrobenzimidazole, dihydro-3H-benzoxazole, phthalazine, [1,8]naphthalidine, [1,5]naphthalidine, 1H-pyrrolo[3,2-c]pyridine, 1H-pyrrolo[2,3-

c]pyridine, 1H-pyrazolo[4,3-b]pyridine, 1H-pyrazolo[4,3-c]pyridine, 1H-pyrazolo[3,4-c]pyridine, 1H-pyrazolo[3,4-b]pyridine, [1,2,4]triazolo[4,3-a]pyridine, thieno[3,2-c]pyridine, thieno[3,2-b]pyridine, 1H-thieno[3,2-c]pyrazole, benzo[d]isoxazole, benzo[c]isoxazole, indolizine, 1,3-dihydroindole, 1H-pyrazolo[3,4-d]thiazole, 2H-isoindole, [1,2,4]triazolo[1,5-a]pyrimidine, 1H-pyrazolo[3,4-b]pyrazine, 1H-imidazo[4,5-b]pyrazine, 7H-purine, or 4H-chromene (the aforementioned residue may be substituted with one of Xa or two or more of the same or different Xa, when AR' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when AR' contains amino group, the amino group may be protected with Rp<sup>2</sup>).

(132) The compound according to (129) or (130) mentioned above, wherein, in the formula (III), AR' is naphthalen-2-yl group, naphthalen-1-yl group, benzofuran-5-yl group, benzofuran-4-yl group, benzofuran-2-yl group, benzo[b]thiophen-5-yl group, benzo[b]thiophen-4-yl group, benzo[b]thiophen-2-yl group, indol-5-yl group, indol-4-yl group, indol-6-yl group, benzothiazol-6-yl group, benzothiazol-7-yl group, benzothiazol-5-yl group, benzothiazol-4-yl group, dihydro-3H-benzothiazol-6-yl group, dihydro-3H-benzothiazol-7-yl group, dihydro-3H-benzothiazol-5-yl group, dihydro-3H-benzothiazol-4-yl group, quinolin-6-yl group, quinolin-3-yl group, quinolin-5-yl group, quinolin-7-yl group, dihydro-1H-quinolin-6-yl group, dihydro-1H-quinolin-5-yl group, benzo[d]isothiazol-5-yl group, benzo[d]isothiazol-4-yl group, benzo[d]isothiazol-6-yl group, benzo[d]isothiazol-7-yl group, 1H-indazol-5-yl group, 1H-indazol-4-yl group, 1H-indazol-6-yl group, benzo[c]isothiazol-5-yl group, benzo[c]isothiazol-4-yl group, benzo[c]isothiazol-6-yl group, benzo[c]isothiazol-7-yl group, 2H-indazol-5-yl group, 2H-indazol-4-yl group, 2H-indazol-6-yl group, imidazo[1,2-a]pyridin-6-yl group, imidazo[1,2-a]pyridin-7-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1H-pyrrolo[2,3-b]pyridin-4-yl group, isoquinolin-6-yl group, isoquinolin-3-yl group, isoquinolin-5-yl group, isoquinolin-7-yl group, dihydro-2H-isoquinolin-6-yl group, dihydro-2H-isoquinolin-5-yl group, cinnolin-6-yl group,

cinnolin-5-yl group, quinazolin-6-yl group, quinazolin-7-yl group, quinazolin-5-yl group, quinoxalin-2-yl group, quinoxalin-6-yl group, quinoxalin-5-yl group, 1H-benzimidazol-5-yl group, 1H-benzimidazol-4-yl group, benzoxazol-5-yl group, benzoxazol-6-yl group, benzoxazol-4-yl group, benzoxazol-7-yl group, 1H-pyrrolo[3,2-b]pyridin-5-yl group, 1H-pyrrolo[3,2-b]pyridin-6-yl group, benzo[1,2,5]thiadiazol-5-yl group, benzo[1,2,5]thiadiazol-4-yl group, 1H-benzotriazol-5-yl group, 1H-benzotriazol-4-yl group, 1,3-dihydropyrrolo[2,3-b]pyridin-5-yl group, 1,3-dihydropyrrolo[2,3-b]pyridin-4-yl group, 1,3-dihydrobenzimidazol-5-yl group, 1,3-dihydrobenzimidazol-4-yl group, dihydro-3H-benzoxazol-6-yl group, dihydro-3H-benzoxazol-7-yl group, dihydro-3H-benzoxazol-5-yl group, dihydro-3H-benzoxazol-4-yl group, phthalazin-6-yl group, phthalazin-5-yl group, [1,8]naphthalidin-3-yl group, [1,8]naphthalidin-4-yl group, [1,5]naphthalidin-3-yl group, [1,5]naphthalidin-4-yl group, 1H-pyrrolo[3,2-c]pyridin-6-yl group, 1H-pyrrolo[3,2-c]pyridin-4-yl group, 1H-pyrrolo[2,3-c]pyridin-5-yl group, 1H-pyrrolo[2,3-c]pyridin-4-yl group, 1H-pyrazolo[4,3-b]pyridin-5-yl group, 1H-pyrazolo[4,3-b]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-4-yl group, 1H-pyrazolo[3,4-c]pyridin-5-yl group, 1H-pyrazolo[3,4-c]pyridin-4-yl group, 1H-pyrazolo[3,4-b]pyridin-5-yl group, 1H-pyrazolo[3,4-b]pyridin-4-yl group, [1,2,4]triazolo[4,3-a]pyridin-6-yl group, [1,2,4]triazolo[4,3-a]pyridin-7-yl group, thieno[3,2-c]pyridin-2-yl group, thieno[3,2-c]pyridin-3-yl group, thieno[3,2-c]pyridin-6-yl group, thieno[3,2-b]pyridin-2-yl group, thieno[3,2-b]pyridin-3-yl group, thieno[3,2-b]pyridin-5-yl group, thieno[3,2-b]pyridin-6-yl group, 1H-thieno[3,2-c]pyrazol-5-yl group, 1H-thieno[3,2-c]pyrazol-4-yl group, benzo[d]isoxazol-5-yl group, benzo[d]isoxazol-4-yl group, benzo[d]isoxazol-6-yl group, benzo[d]isoxazol-7-yl group, benzo[c]isoxazol-5-yl group, benzo[c]isoxazol-4-yl group, benzo[c]isoxazol-6-yl group, benzo[c]isoxazol-7-yl group, indolizin-7-yl group, indolizin-6-yl group, indolizine-8-yl group, 1,3-dihydroindol-5-

yl group, 1,3-dihydroindol-4-yl group, 1,3-dihydroindol-6-yl group, 1H-pyrazolo[3,4-d]thiazol-5-yl group, 2H-isoindol-5-yl group, 2H-isoindol-4-yl group, [1,2,4]triazolo[1,5-a]pyrimidin-6-yl group, 1H-pyrazolo[3,4-b]pyrazin-5-yl group, 1H-imidazo[4,5-b]pyrazin-5-yl group, 7H-purin-2-yl group, 4H-chromen-6-yl group, or 4H-chromen-5-yl group (the aforementioned groups may be substituted with one of Xa or two or more of the same or different Xa, when AR' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when AR' contains amino group, the amino group may be protected with Rp<sup>2</sup>).

(133) The compound according to (129) or (130) mentioned above, wherein, in the formula (III), AR' is a residue of naphthalene, benzofuran, benzo[b]thiophene, indole, benzothiazole, dihydro-3H-benzothiazole, quinoline, dihydro-1H-quinoline, benzo[d]isothiazole, 1H-indazole, benzo[c]isothiazole, 2H-indazole, imidazo[1,2-a]pyridine, 1H-pyrrolo[2,3-b]pyridine, isoquinoline, or dihydro-2H-isoquinoline (the aforementioned residue may be substituted with one of Xa or two or more of the same or different Xa, when AR' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when AR' contains amino group, the amino group may be protected with Rp<sup>2</sup>).

(134) The compound according to (129) or (130) mentioned above, wherein, in the formula (III), AR' is a residue of cinnoline, quinazoline, quinoxaline, 1H-benzimidazole, benzoxazole, 1H-pyrrolo[3,2-b]pyridine, benzo[1,2,5]thiadiazole, 1H-benzotriazole, 1,3-dihydropyrrolo[2,3-b]pyridine, 1,3-dihydrobenzimidazole, dihydro-3H-benzoxazole, phthalazine, [1,8]naphthalidine, [1,5]naphthalidine, 1H-pyrrolo[3,2-c]pyridine, 1H-pyrrolo[2,3-c]pyridine, 1H-pyrazolo[4,3-b]pyridine, 1H-pyrazolo[4,3-c]pyridine, 1H-pyrazolo[3,4-c]pyridine, 1H-pyrazolo[3,4-b]pyridine, [1,2,4]triazolo[4,3-a]pyridine, thieno[3,2-c]pyridine, thieno[3,2-b]pyridine, 1H-thieno[3,2-c]pyrazole, benzo[d]isoxazole, benzo[c]isoxazole, indolizine, 1,3-dihydroindole, 1H-pyrazolo[3,4-d]thiazole, 2H-isoindole, [1,2,4]triazolo[1,5-

alpyrimidine, 1H-pyrazolo[3,4-b]pyrazine, 1H-imidazo[4,5-b]pyrazine, 7H-purine, or 4H-chromene (the aforementioned residue may be substituted with one of Xa or two or more of the same or different Xa, when AR' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when AR' contains amino group, the amino group may be protected with Rp<sup>2</sup>).

(135) The compound according to any one of (129) to (134) mentioned above, wherein, in the formula (III), Rs' is -O-Rx'.

(136) The compound according to any one of (129) to (135) mentioned above, wherein, in the formula (III), Rs' is -D-Rx' or -N(Ry')(Rz'), D is a single bind, oxygen atom, sulfur atom, -S(O)-, -S(O)<sub>2</sub>-, or -C(O)-, Rx' is a linear or branched saturated alkyl group having 3 to 8 carbon atoms, or Ra, Rb, or Rc, k in Ra is 0 or an integer of 1 to 3, R<sup>1</sup> is a saturated cyclic alkyl group having 3 to 7 carbon atoms or a condensed saturated cyclic alkyl group having 6 to 8 carbon atoms, R<sup>1</sup> may be substituted with one of lower alkyl group having 1 to 4 carbon atoms or two or more of the same or different lower alkyl groups having 1 to 4 carbon atoms, Q in Rb is phenyl group, thienyl group, furyl group, pyrrolyl group, pyridyl group, oxazolyl group, isoxazolyl group, thiazolyl group, isothiazolyl group, imidazolyl group, pyrazolyl group, oxadiazolyl group, thiadiazolyl group, triazolyl group, tetrazolyl group, naphthyl group, tetrahydronaphthyl group, indanyl group, indenyl group, quinolyl group, isoquinolyl group, indolyl group, benzofuryl group, benzothienyl group, benzimidazolyl group, benzoxazolyl group, benzothiazolyl group, indazolyl group, 4H-chromenyl group, dihydrobenzodioxyl group, benzoisoxazolyl group, pyrrolopyridinyl group, pyrazolopyridinyl group, triazolopyridinyl group, thienopyridinyl group, thienopyrazolyl group, 1,3-dihydrobenzimidazole group, dihydro-3H-benzoxazole group, or dihydro-3H-benzothiazole group, which binds to A<sup>2</sup> at an arbitrary position on the ring, A<sup>1</sup> is a single bind or an alkylene (a) having 1 to 3 carbon atoms, the alkylene (a) may be substituted with a lower alkyl group



having 1 to 4 carbon atoms or phenyl group may be substituted with, A<sup>2</sup> is a single bond, oxygen atom, sulfur atom, -S(O)-, -S(O)<sub>2</sub>-, or -N(R<sup>4</sup>)- (provided that when A<sup>2</sup> is oxygen atom, sulfur atom, -S(O)-, -S(O)<sub>2</sub>-, or -N(R<sup>4</sup>)-, A<sup>1</sup> is ethylene or trimethylene), R<sup>2</sup> and R<sup>3</sup> independently represent hydrogen atom, a linear or branched saturated alkyl group having 1 to 4 carbon atoms, oxo group, thioxo group, fluorine atom, chlorine atom, bromine atom, trifluoromethyl group, -OR<sup>5</sup>, -N(R<sup>6</sup>)(R<sup>6'</sup>), -NHCOR<sup>7</sup>, -NHSO<sub>2</sub>R<sup>8</sup>, or -A<sup>6</sup>-Qa, or they binds to each other to form methylenedioxy group, Qa is phenyl group, pyridyl group, oxazolyl group, isoxazolyl group, thiazolyl group, isothiazolyl group, imidazolyl group, pyrazolyl group, oxadiazolyl group, thiadiazolyl group, triazolyl group, tetrazolyl group, naphthyl group, indanyl group, indenyl group, quinolyl group, isoquinolyl group, indolyl group, benzofuryl group, benzothienyl group, benzimidazolyl group, benzoxazolyl group, benzothiazolyl group, or indazolyl group, which may be substituted with one of T<sup>1</sup> or two or more of the same or different T<sup>1</sup>, and binds to A<sup>6</sup> at an arbitrary position on the ring, R<sup>4</sup> and R<sup>6</sup> independently represent hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms, R<sup>5</sup> and R<sup>7</sup> independently represent hydrogen atom, a lower alkyl group having 1 to 4 carbon atoms, or -A<sup>6</sup>-Qa, R<sup>8</sup> is a lower alkyl group having 1 to 4 carbon atoms, R<sup>6'</sup> has the same meaning as R<sup>6</sup>, or binds to R<sup>6</sup> to form a 3- to 6-membered ring together with the nitrogen atom to which they bind to form a saturated nitrogen-containing alkyl group or morpholino group, p in Rc is an integer of 2 to 4, A<sup>4</sup> is a single bond or methylene or ethylene, A<sup>5</sup> is -C(O)-, -C(S)-, or -S(O)<sub>2</sub>-, Rd is hydrogen atom, an alkyl group having 1 to 8 carbon atoms, or Qa, Re is an alkyl group having 1 to 8 carbon atoms, -A<sup>6</sup>-Qa, -(CH<sub>2</sub>)<sub>i</sub>R<sup>14</sup>, -OR<sup>28</sup>, -SR<sup>28</sup>, or -N(R<sup>29</sup>)(R<sup>30</sup>), i is an integer of 1 to 3, R<sup>14</sup> is hydroxyl group, an alkoxy group having 1 to 4 carbon atoms, carboxyl group, or an N,N-dialkylcarbamoyl group having 1 to 4 carbon atoms, R<sup>28</sup> is an alkyl group having 1

to 8 carbon atoms or  $-A^6-Qa$ ,  $R^{29}$  is an alkyl group having 1 to 8 carbon atoms, an alkoxy carbonyl group having 1 to 4 carbon atoms, or  $-A^6-Qa$ ,  $R^{30}$  is hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms, or binds to  $R^{29}$  to form a 3- to 6-membered ring together with the nitrogen atom to which they bind to form a saturated nitrogen-containing alkyl group or morpholino group,  $Rz'$  has the same meaning as  $Rx'$ , or represents  $-A^5-Re$ ,  $Ry'$  is hydrogen atom, an alkyl group having 1 to 8 carbon atoms, or  $-A^6-Qp$ , or binds to  $Rz'$  to form a saturated or unsaturated nitrogen-containing cyclic substituent having 3 to 7 atoms, when  $-D-Rx'$  or  $-N(Ry')(Rz')$  contains hydroxyl group, the hydroxyl group may be protected with  $Rp^1$ , and when  $-D-Rx'$  or  $-N(Ry')(Rz')$  contains amino group, the amino group may be protected with  $Rp^2$ .

(137) The compound according to any one of (129) to (136) mentioned above, wherein, in the formula (III),  $Rs'$  is  $-N(Ry')(Rz')$ .

(138) The compound according to any one of (129) to (136) mentioned above, wherein, in the formula (III),  $Rs'$  is  $-D-Rx'$ ,  $D$  is sulfur atom,  $-S(O)-$ ,  $-S(O)_2-$ , or  $-C(O)-$ .

(139) The compound according to (129) mentioned above, wherein, in the formula (III),  $AR'$  binds at the position of  $C^{2'}$  in the aromatic ring ( $E'$ ), and  $Rs'$  binds to one of the ring-constituting carbon atoms  $C^{3'}$ ,  $C^{4'}$ , and  $C^{5'}$ .

(140) The compound according to (129) mentioned above, wherein, in the formula (III),  $AR'$  binds to  $C^{2'}$ ,  $Rs'$  binds to any one of the atoms  $C^{3'}$ ,  $C^{4'}$ , and  $C^{5'}$ , a ring-constituting carbon atom to which  $Rs'$  does not bind among  $C^{3'}$ ,  $C^{4'}$ , and  $C^{5'}$  may be replaced with  $V'$ ,

$V'$  is nitrogen atom, or carbon atom substituted with  $Zx'$ ,  $Zx'$  is one kind of group selected from the group consisting of fluorine atom, chlorine atom, bromine atom, nitro group, methyl group, hydroxyl group, methoxy group, amino group, N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino

group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, and N,N-dimethylsulfamoylamino group, provided that when Zx' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when Zx' contains amino group, the amino group may be protected with Rp<sup>2</sup>,

Rs' is -D-Rx' or -N(Ry')(Rz'), D is oxygen atom or sulfur atom, Rx' is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-cyclopentylethyl group, or 2-cyclohexylethyl group, or Rb or Rc, Q in Rb is phenyl group, thienyl group, furyl group, pyridyl group, oxazolyl group, naphthyl group, tetrahydronaphthyl group, indanyl group, indolyl group, or dihydrobenzodioxyl group, A<sup>2</sup> is a single bind, oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)- (provided that when A<sup>2</sup> is oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)-, A<sup>1</sup> is ethylene), R<sup>2</sup> and R<sup>3</sup> independently represent hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, dimethylamino group, acetylamino group, or methylsulfonylamino group (provided that when Q is phenyl group, A<sup>1</sup> is a single bind or unsubstituted methylene, and A<sup>2</sup> is a single bind, one of R<sup>2</sup> and R<sup>3</sup> is a substituent other than hydrogen atom), p in Rc is an integer of 2 or 3, A<sup>4</sup> is a single bind or methylene, A<sup>5</sup> is -C(O)-, -C(S)-, or -S(O)<sub>2</sub>-, Rd is hydrogen atom, or methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group, cyclopropylmethyl group, cyclopentyl group, cyclopentylmethyl group, cyclohexyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, or pyridin-4-yl group, Re is methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl

group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, phenylmethyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, furan-2-yl group, furan-3-yl group, thiophen-2-yl group, thiophen-3-yl group, methoxy group, ethoxy group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenyloxy group, 4-methylphenyloxy group, 4-chlorophenyloxy group, 4-fluorophenyloxy group, thiomethoxy group, amino group, N-methylamino group, N,N-dimethylamino group, N-ethylamino group, N,N-diethylamino group, N-propylamino group, N-isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group, N-(4-fluorophenyl)amino group, N-(pyridin-2-yl)amino group, N-(pyridin-3-yl)amino group, N-(pyridin-4-yl)amino group, N-(furan-2-yl)amino group, N-(furan-3-yl)amino group, N-(thiophen-2-yl)amino group, N-(thiophen-3-yl)amino group, pyrrolidino group, piperidino group, morpholino group, methyloxycarbonylamino group, or ethyloxycarbonylamino group, R<sub>z</sub>' is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-

2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-  
 dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-  
 (2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl  
 group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-  
 chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl  
 group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-  
 dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl  
 group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-  
 chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl  
 group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-  
 difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-  
 dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-  
 dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-  
 dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-  
 (trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-  
 (trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-  
 methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-  
 methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-  
 methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl  
 group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-  
 chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-  
 (trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-  
 (trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group,  
 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-  
 chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl  
 group, 2-(N-phenyl-N-methylamino)ethyl group, 2-(N-ethyl-N-phenylamino)ethyl  
 group, isobutyryl group, isopropylthiocarbonyl group, isopropylsulfonyl group,

valeryl group, butylthiocarbonyl group, isovaleryl group, isobutylthiocarbonyl group, pivaloyl group, t-butylthiocarbonyl group, cyclopropylcarbonyl group, cyclopropylthiocarbonyl group, cyclopentylcarbonyl group, cyclopentylthiocarbonyl group, cyclohexylcarbonyl group, cyclohexylthiocarbonyl group, cyclopentylmethylcarbonyl group, cyclopentylmethylthiocarbonyl group, cyclohexylmethylcarbonyl group, cyclohexylmethylthiocarbonyl group, benzoyl group, thiobenzoyl group, phenylsulfonyl group, 4-methylphenylcarbonyl group, 4-methylphenylthiocarbonyl group, 4-methylphenylsulfonyl group, 4-chlorophenylcarbonyl group, 4-chlorophenylthiocarbonyl group, 4-fluorophenylcarbonyl group, 4-fluorophenylthiocarbonyl group, isopropoxy carbonyl group, N-isopropyl carbamoyl group, N-isopropylthiocarbamoyl group, butyloxy carbonyl group, N-butyl carbamoyl group, N-butylthiocarbamoyl group, isobutyloxy carbonyl group, N-isobutyl carbamoyl group, N-isobutylthiocarbamoyl group, t-butyloxy carbonyl group, N-t-butyl carbamoyl group, N-t-butylthiocarbamoyl group, cyclopropyloxy carbonyl group, N-cyclopropyl carbamoyl group, N-cyclopropylthiocarbamoyl group, cyclopentyloxy carbonyl group, N-cyclopentyl carbamoyl group, N-cyclopentylthiocarbamoyl group, cyclohexyloxy carbonyl group, N-cyclohexyl carbamoyl group, N-cyclohexylthiocarbamoyl group, cyclopentylmethyloxy carbonyl group, cyclohexylmethyloxy carbonyl group, phenyloxy carbonyl group, N-phenyl carbamoyl group, N-phenylthiocarbamoyl group, 4-methylphenyloxy carbonyl group, N-(4-methylphenyl) carbamoyl group, N-(4-methylphenyl)thiocarbamoyl group, 4-chlorophenyloxy carbonyl group, N-(4-chlorophenyl) carbamoyl group, N-(4-chlorophenyl)thiocarbamoyl group, 4-fluorophenyloxy carbonyl group, N-(4-fluorophenyl) carbamoyl group, N-(4-fluorophenyl)thiocarbamoyl group, (pyrrolidino-1-yl) carbonyl group, (piperidino-1-yl) carbonyl group, or (morpholino-4-yl) carbonyl group, Ry' is hydrogen atom, methyl

group, ethyl group or isobutyl group, or binds to R<sub>z</sub>' to form pyrrolidino group, piperidino group, piperazino group, morpholino group, pyrrol-1-yl group, imidazol-1-yl group or pyrazol-1-yl group together with the nitrogen atom, provided that when -D·R<sub>x</sub>' or -N(R<sub>y</sub>') (R<sub>z</sub>') contains hydroxyl group, the hydroxyl group may be protected with R<sub>p</sub><sup>1</sup>, and when -D·R<sub>x</sub>' or -N(R<sub>y</sub>') (R<sub>z</sub>') contains amino group, the amino group may be protected with R<sub>p</sub><sup>2</sup>,

AR' is naphthalen-2-yl group, naphthalen-1-yl group, benzofuran-5-yl group, benzofuran-4-yl group, benzofuran-2-yl group, benzo[b]thiophen-5-yl group, benzo[b]thiophen-4-yl group, benzo[b]thiophen-2-yl group, indol-5-yl group, indol-4-yl group, indol-6-yl group, benzothiazol-6-yl group, benzothiazol-7-yl group, benzothiazol-5-yl group, benzothiazol-4-yl group, dihydro-3H-benzothiazol-6-yl group, dihydro-3H-benzothiazol-7-yl group, dihydro-3H-benzothiazol-5-yl group, dihydro-3H-benzothiazol-4-yl group, quinolin-6-yl group, quinolin-3-yl group, quinolin-5-yl group, quinolin-7-yl group, dihydro-1H-quinolin-6-yl group, dihydro-1H-quinolin-5-yl group, benzo[d]isothiazol-5-yl group, benzo[d]isothiazol-4-yl group, benzo[d]isothiazol-6-yl group, benzo[d]isothiazol-7-yl group, 1H-indazol-5-yl group, 1H-indazol-4-yl group, 1H-indazol-6-yl group, benzo[c]isothiazol-5-yl group, benzo[c]isothiazol-4-yl group, benzo[c]isothiazol-6-yl group, benzo[c]isothiazol-7-yl group, 2H-indazol-5-yl group, 2H-indazol-4-yl group, 2H-indazol-6-yl group, imidazo[1,2-a]pyridin-6-yl group, imidazo[1,2-a]pyridin-7-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1H-pyrrolo[2,3-b]pyridin-4-yl group, isoquinolin-6-yl group, isoquinolin-3-yl group, isoquinolin-5-yl group, isoquinolin-7-yl group, dihydro-2H-isoquinolin-6-yl group, dihydro-2H-isoquinolin-5-yl group, cinnolin-6-yl group, cinnolin-5-yl group, quinazolin-6-yl group, quinazolin-7-yl group, quinazolin-5-yl group, quinoxalin-2-yl group, quinoxalin-6-yl group, quinoxalin-5-yl group, 1H-benzimidazol-5-yl group, 1H-benzimidazol-4-yl group, benzoxazol-5-yl group, benzoxazol-6-yl group, benzoxazol-4-yl group, benzoxazol-7-yl group, 1H-

pyrrolo[3,2-b]pyridin-5-yl group, 1H-pyrrolo[3,2-b]pyridin-6-yl group,  
 benzo[1,2,5]thiadiazol-5-yl group, benzo[1,2,5]thiadiazol-4-yl group, 1H-  
 benzotriazol-5-yl group, 1H-benzotriazol-4-yl group, 1,3-dihydropyrrolo[2,3-  
 b]pyridin-5-yl group, 1,3-dihydropyrrolo[2,3-b]pyridin-4-yl group, 1,3-  
 dihydrobenzimidazol-5-yl group, 1,3-dihydrobenzimidazol-4-yl group, dihydro-3H-  
 benzoxazol-6-yl group, dihydro-3H-benzoxazol-7-yl group, dihydro-3H-benzoxazol-5-  
 yl group, dihydro-3H-benzoxazol-4-yl group, phthalazin-6-yl group, phthalazin-5-yl  
 group, [1,8]naphthalidin-3-yl group, [1,8]naphthalidin-4-yl group,  
 [1,5]naphthalidin-3-yl group, [1,5]naphthalidin-4-yl group, 1H-pyrrolo[3,2-  
 c]pyridin-6-yl group, 1H-pyrrolo[3,2-c]pyridin-4-yl group, 1H-pyrrolo[2,3-c]pyridin-  
 5-yl group, 1H-pyrrolo[2,3-c]pyridin-4-yl group, 1H-pyrazolo[4,3-b]pyridin-5-yl  
 group, 1H-pyrazolo[4,3-b]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-6-yl group,  
 1H-pyrazolo[4,3-c]pyridin-4-yl group, 1H-pyrazolo[3,4-c]pyridin-5-yl group, 1H-  
 pyrazolo[3,4-c]pyridin-4-yl group, 1H-pyrazolo[3,4-b]pyridin-5-yl group, 1H-  
 pyrazolo[3,4-b]pyridin-4-yl group, [1,2,4]triazolo[4,3-a]pyridin-6-yl group,  
 [1,2,4]triazolo[4,3-a]pyridin-7-yl group, thieno[3,2-c]pyridin-2-yl group, thieno[3,2-  
 c]pyridin-3-yl group, thieno[3,2-c]pyridin-6-yl group, thieno[3,2-b]pyridin-2-yl group,  
 thieno[3,2-b]pyridin-3-yl group, thieno[3,2-b]pyridin-5-yl group, thieno[3,2-  
 b]pyridin-6-yl group, 1H-thieno[3,2-c]pyrazol-5-yl group, 1H-thieno[3,2-c]pyrazol-4-  
 yl group, benzo[d]isoxazol-5-yl group, benzo[d]isoxazol-4-yl group, benzo[d]isoxazol-  
 6-yl group, benzo[d]isoxazol-7-yl group, benzo[c]isoxazol-5-yl group,  
 benzo[c]isoxazol-4-yl group, benzo[c]isoxazol-6-yl group, benzo[c]isoxazol-7-yl group,  
 indolizin-7-yl group, indolizin-6-yl group, indolizine-8-yl group, 1,3-dihydroindol-5-  
 yl group, 1,3-dihydroindol-4-yl group, 1,3-dihydroindol-6-yl group, 1H-pyrazolo[3,4-  
 d]thiazol-5-yl group, 2H-isoindol-5-yl group, 2H-isoindol-4-yl group,  
 [1,2,4]triazolo[1,5-a]pyrimidin-6-yl group, 1H-pyrazolo[3,4-b]pyrazin-5-yl group,  
 1H-imidazo[4,5-b]pyrazin-5-yl group, 7H-purin-2-yl group, 4H-chromen-6-yl group,



or 4H-chromen-5-yl group (the aforementioned groups may be substituted with one of Xa or two or more of the same or different Xa), and Xa is oxo group, thioxo group, fluorine atom, chlorine atom, trifluoromethyl group, methyl group, ethyl group, propyl group, 2-hydroxyethyl group, carboxymethyl group, 2-carboxyethyl group, N,N-dimethylcarbamoylmethyl group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, carboxymethyloxy group, 2-carboxyethyloxy group, N,N-dimethylcarbamoylmethyloxy group, amino group, methylamino group, dimethylamino group, 2-hydroxyethylamino group, carbamoylamino group, acetylamino group, furan-2-carboxyamino group, 2-hydroxyacetylamino group, 2-aminoacetylamino group, methylsulfonylamino group, (N,N-dimethylsulfamoyl)amino group, methanesulfonyl group, sulfamoyl group, N-methylsulfamoyl group, N,N-dimethylsulfamoyl group, carboxyl group, acetyl group, carbamoyl group, or N,N-dimethylcarbamoyl group, provided that when AR' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when AR' contains amino group, the amino group may be protected with Rp<sup>2</sup>.

(141) The compound according to (139) or (140) mentioned above, wherein, in the formula (III), Rs' is -O-Rx', and all of C<sup>2'</sup>, C<sup>3'</sup>, C<sup>4'</sup>, C<sup>5'</sup> and C<sup>6'</sup> in the aromatic ring (E') are not replaced with V'.

(142) The compound according to (131) mentioned above, wherein, in the formula (III), AR' binds at the position of C<sup>2'</sup> in the aromatic ring (E'), Rs' binds to one of the ring-constituting carbon atoms C<sup>3'</sup>, C<sup>4'</sup>, and C<sup>5'</sup>, Rs' is -O-Rx', and all of C<sup>2'</sup>, C<sup>3'</sup>, C<sup>4'</sup>, C<sup>5'</sup> and C<sup>6'</sup> in the aromatic ring (E') are not replaced with V.

(143) The compound according to (132) mentioned above, wherein, in the formula (III), AR' binds at the position of C<sup>2'</sup> in the aromatic ring (E'), Rs' binds to one of the ring-constituting carbon atoms C<sup>3'</sup>, C<sup>4'</sup>, and C<sup>5'</sup>, Rs' is -O-Rx', and all of C<sup>2'</sup>, C<sup>3'</sup>, C<sup>4'</sup>, C<sup>5'</sup> and C<sup>6'</sup> in the aromatic ring (E') are not replaced with V.

(144) The compound according to any one of (139) to (143) mentioned above,

wherein, in the formula (III), Rs' binds to C<sup>3'</sup>.

(145) The compound according to (129) mentioned above, wherein, in the formula (III), C<sup>2'</sup> is carbon atom to which AR' binds, C<sup>3'</sup> is carbon atom to which Rs' binds, C<sup>4'</sup> may be replaced with V', C<sup>5'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms,

V' is nitrogen atom, or carbon atom substituted with Zx', Zx' is fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, or N,N-dimethylamino group, provided that when Zx' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when Zx' contains amino group, the amino group may be protected with Rp<sup>2</sup>,

Rs' is -O-Rx', Rx' is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-

chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, or 2-(N-ethyl-N-phenylamino)ethyl group, and

AR' is naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-

indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group, 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, or benzoxazol-5-yl group (the aforementioned groups may be substituted with one of Xa or two or more of the same or different Xa), provided that when AR' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when AR' contains amino group, the amino group may be protected with Rp<sup>2</sup>.

(146) The compound according to (139) to (143) mentioned above, wherein, in the formula (III), Rs' binds to C<sup>4'</sup>.

(147) The compound according to (129) mentioned above, wherein, in the formula (III), C<sup>2'</sup> is carbon atom to which AR' binds, C<sup>4'</sup> is carbon atom to which Rs' binds, C<sup>5'</sup> may be replaced with V', C<sup>3'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms,

V' is nitrogen atom, or carbon atom substituted with Zx', Zx' is fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, or N,N-dimethylamino group, provided that when Zx' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when Zx' contains amino group, the amino group may be protected with Rp<sup>2</sup>,

Rs' is -O-Rx', Rx' is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl

group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, or 2-(N-ethyl-N-phenylamino)ethyl group, and

AR' is naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-

methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group, 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, or benzoxazol-5-yl group, provided that when AR' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when AR' contains amino group, the amino group may be protected with Rp<sup>2</sup>.

(148) The compound according to any one of (139) to (143) mentioned above, wherein, in the formula (III), Rs' binds to C<sup>5'</sup>.

(149) The compound according to (129) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to the atom C<sup>5'</sup> or C<sup>6'</sup> in the aromatic ring (E').

(150) The compound according to (149) mentioned above, wherein, in the formula (III), Rs' is -O-Rx', and all of C<sup>2'</sup>, C<sup>3'</sup>, C<sup>4'</sup>, C<sup>5'</sup> and C<sup>6'</sup> in the aromatic ring (E') are not replaced with V.

(151) The compound according to (131) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to the atom C<sup>5'</sup> or C<sup>6'</sup> in the aromatic ring (E'), Rs' is -O-Rx', and all of C<sup>2'</sup>, C<sup>3'</sup>, C<sup>4'</sup>, C<sup>5'</sup> and C<sup>6'</sup> in the aromatic ring (E') are not replaced with V.

(152) The compound according to (132) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to the atom C<sup>5'</sup> or C<sup>6'</sup> in the aromatic ring (E'), Rs' is -O-Rx', and all of C<sup>2'</sup>, C<sup>3'</sup>, C<sup>4'</sup>, C<sup>5'</sup> and C<sup>6'</sup> in the aromatic ring (E') are not replaced with V.

(153) The compound according to any one of (149) to (152) mentioned above, wherein, in the formula (III), Rs' binds to C<sup>5'</sup>.

(154) The compound according to (129) mentioned above, wherein, in the formula (III), C<sup>3'</sup> is carbon atom to which AR' binds, C<sup>5'</sup> is carbon atom to which Rs' binds, C<sup>2'</sup>, C<sup>4'</sup>, and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms,

Rs' is -O-Rx', Rx' is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group,



5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group,  
 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group,  
 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group,  
 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl  
 group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl  
 group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-  
 chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl  
 group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-  
 methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-  
 dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl  
 group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-  
 chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl  
 group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-  
 difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-  
 dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-  
 dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-  
 dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-  
 (trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-  
 (trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-  
 methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-  
 methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-  
 methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl  
 group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-  
 chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-  
 (trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-  
 (trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group,  
 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-

chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, or 2-(N-ethyl-N-phenylamino)ethyl group, and

AR' is naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group, 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-

5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, or benzoxazol-5-yl group, provided that when AR' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when AR' contains amino group, the amino group may be protected with Rp<sup>2</sup>.

(155) The compound according to any one of (149) to (152) mentioned above, wherein, in the formula (III), Rs' binds to C<sup>6'</sup>.

(156) The compound according to (129) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), and C<sup>6'</sup> is V'.

(157) The compound according to (131) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>6'</sup> is carbon atom substituted with Zx, C<sup>2'</sup> and C<sup>5'</sup> are unsubstituted carbon atoms, and Rs' is -O-Rx'.

(158) The compound according to (132) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>6'</sup> is carbon atom substituted with Zx, C<sup>2'</sup> and C<sup>5'</sup> are unsubstituted ring-constituting carbon atoms, and Rs' is -O-Rx'.

(159) The compound according to (129) mentioned above, wherein, in the formula (III), C<sup>3'</sup> is carbon atom to which AR' binds, C<sup>4'</sup> is a carbon atom to which Rs' binds, C<sup>6'</sup> is carbon atom substituted with Zx', C<sup>2'</sup> and C<sup>5'</sup> are unsubstituted ring-constituting carbon atoms,

Zx' is fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, or N,N-dimethylamino group, provided that when Zx' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when Zx' contains amino group, the amino group may be protected with Rp<sup>2</sup>,

Rs' is -O-Rx', Rx' is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-

(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, or 2-(N-ethyl-N-phenylamino)ethyl group, and

AR' is naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-

indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group, 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, or benzoxazol-5-yl group, provided that when Ar' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when AR' contains amino group, the amino group may be protected with Rp<sup>2</sup>.

(160) The compound according to (129) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5'</sup> is nitrogen atom, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms, and Rs' is -O-Rx'.

(161) The compound according to (131) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5'</sup> is nitrogen atom, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon

atoms, and Rs' is -O-Rx'.

(162) The compound according to (132) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5'</sup> is nitrogen atom, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms, and Rs' is -O-Rx'.

(163) The compound according to (129) mentioned above, wherein, in the formula (III), C<sup>3'</sup> is carbon atom to which AR' binds, C<sup>4'</sup> is carbon atom to which Rs' binds, C<sup>5'</sup> is nitrogen atom, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms,

Rs' is -O-Rx', Rx' is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-

difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, or 2-(N-ethyl-N-phenylamino)ethyl group, and

AR' is naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-



5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group, 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, or benzoxazol-5-yl group, provided that when AR' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when AR' contains amino group, the amino group may be protected with Rp<sup>2</sup>.

(164) The compound according to (129) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5'</sup> is a ring-constituting carbon atom substituted with Zx', or an unsubstituted

ring-constituting carbon atom, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms, Rs' is -D-Rx', D is a single bind, sulfur atom, -S(O)-, -S(O)<sub>2</sub>-, or -C(O)-.

(165) The compound according to (131) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5</sup> is a ring-constituting carbon atom substituted with Zx', or an unsubstituted ring-constituting carbon atom, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms, Rs' is -D-Rx', and D is a single bind, sulfur atom, -S(O)-, -S(O)<sub>2</sub>-, or -C(O)-.

(166) The compound according to (132) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5'</sup> is a ring-constituting carbon atom substituted with Zx', or an unsubstituted ring-constituting carbon atom, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms, Rs' is -D-Rx', and D is a single bind, sulfur atom, -S(O)-, -S(O)<sub>2</sub>-, or -C(O)-.

(167) The compound according to (129) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5'</sup> is a ring-constituting carbon atom substituted with Zx', or an unsubstituted ring-constituting carbon atom, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms, and Rs' is -N(Ry')(Rz').

(168) The compound according to (131) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5</sup> is a ring-constituting carbon atom substituted with Zx', or an unsubstituted ring-constituting carbon atom, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms, and Rs' is -N(Ry')(Rz').

(169) The compound according to (132) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5'</sup> is a ring-constituting carbon atom substituted with Zx', or an unsubstituted

ring-constituting carbon atom, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms, and Rs' is -N(Ry')(Rz').

(170) The compound according to (129) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5</sup> is carbon atom substituted with -N(Rn<sup>1</sup>)(Rn<sup>2</sup>) (provided that one of Rn<sup>1</sup> and Rn<sup>2</sup> is a substituent other than hydrogen atom), C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms, and Rs' is -O-Rx'.

(171) The compound according to (131) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5'</sup> is carbon atom substituted with -N(Rn<sup>1</sup>)(Rn<sup>2</sup>) (provided that one of Rn<sup>1</sup> and Rn<sup>2</sup> is a substituent other than hydrogen atom), C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms, and Rs' is -O-Rx'.

(172) The compound according to (123) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5'</sup> is carbon atom substituted with -N(Rn<sup>1</sup>)(Rn<sup>2</sup>) (provided that one of Rn<sup>1</sup> and Rn<sup>2</sup> is a substituent other than hydrogen atom), C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms, and Rs' is -O-Rx'.

(173) The compound according to (129) mentioned above, wherein, in the formula (III), C<sup>3'</sup> is carbon atom to which AR' binds, C<sup>4'</sup> is carbon atom to which Rs' binds, C<sup>5'</sup> is carbon atom substituted with Zx', C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms,

Zx' is N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, or N,N-dimethylsulfamoylamino group, provided that when Zx' contains amino group, the amino group may be protected with Rp<sup>2</sup>,

Rs' is -O-Rx', Rx' is butyl group, isobutyl group, 2-ethylbutyl group,

cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl

group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, or 2-(N-ethyl-N-phenylamino)ethyl group, and

AR' is naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group, 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-

dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, or benzoxazol-5-yl group, provided that when AR' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when AR' contains amino group, the amino group may be protected with Rp<sup>2</sup>.

(174) The compound according to (129) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5'</sup> is a ring-constituting carbon atom substituted with Zx', or an unsubstituted ring-constituting carbon atom, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms, Rs' is -D-Rx', and Rx' has the same meaning as Rc, provided that when Rc contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when Rc contains amino group, the amino group may be protected with Rp<sup>2</sup>.

(175) The compound according to (131) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5'</sup> is a ring-constituting carbon atom substituted with Zx', or an unsubstituted ring-constituting carbon atom, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms, Rs' is -D-Rx', and Rx' has the same meaning as Rc, provided that

when Rc contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when Rc contains amino group, the amino group may be protected with Rp<sup>2</sup>.

(176) The compound according to (132) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5'</sup> is a ring-constituting carbon atom substituted with Zx', or an unsubstituted ring-constituting carbon atom, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms, Rs' is -D-Rx', and Rx' has the same meaning as Rc, provided that when Rc contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when Rc contains amino group, the amino group may be protected with Rp<sup>2</sup>.

(177) The compound according to (129) mentioned above, wherein, in the formula (III), C<sup>3'</sup> is carbon atom to which AR' binds, C<sup>4'</sup> is a carbon atom to which Rs' binds, C<sup>5'</sup> is a ring-constituting carbon atom substituted with Zx', or an unsubstituted ring-constituting carbon atom, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms,

Zx' is fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, or N,N-dimethylamino group, provided that when Zx' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when Zx' contains amino group, the amino group may be protected with Rp<sup>2</sup>,

Rs' is -O-Rx', Rx' has the same meaning as Rc, provided that when Rc contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, p in Rc is an integer of 2, A<sup>4</sup> is a single bond or methylene, A<sup>5</sup> is -C(O)-, -C(S)-, or -S(O)<sub>2</sub>-, Rd is methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, or 4-fluorophenylmethyl group, Re is isopropyl group, butyl group, isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group,

4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenyloxy group, 4-methylphenyloxy group, 4-chlorophenyloxy group, 4-fluorophenyloxy group, N-propylamino group, N-isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group, N-(4-fluorophenyl)amino group, pyrrolidino group, piperidino group, or morpholino group, and

AR' is naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-



indol-5-yl group, benzothiazol-6-yl group, 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, or benzoxazol-5-yl group, provided that when AR' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when AR' contains amino group, the amino group may be protected with Rp<sup>2</sup>.

(178) The compound according to (129) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup>, Rs' binds to any one of the atoms C<sup>4'</sup>, C<sup>5'</sup>, and C<sup>6'</sup>, a ring-constituting carbon atom to which Rs' does not bind among C<sup>4'</sup>, C<sup>5'</sup>, and C<sup>6'</sup> may be replaced with V',

V' is nitrogen atom, or carbon atom substituted with Zx', Zx' is fluorine atom, chlorine atom, bromine atom, nitro group, methyl group, hydroxyl group, methoxy group, amino group, N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, or N,N-dimethylsulfamoylamino group, provided that when Zx'

contains hydroxyl group, the hydroxyl group may be protected with  $Rp^1$ , and when  $Zx'$  contains amino group, the amino group may be protected with  $Rp^2$ ,

$Rs'$  is  $-D-Rx'$  or  $-N(Ry')(Rz')$ ,  $D$  is oxygen atom or sulfur atom,  $Rx'$  is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-cyclopentylethyl group, or 2-cyclohexylethyl group, or  $Rb$  or  $Rc$ ,  $Q$  in  $Rb$  is phenyl group, thienyl group, furyl group, pyridyl group, oxazolyl group, naphthyl group, tetrahydronaphthyl group, indanyl group, indolyl group, or dihydrobenzodioxyl group,  $A^2$  is a single bind, oxygen atom, sulfur atom,  $-N(methyl)-$ , or  $-N(ethyl)-$  (provided that when  $A^2$  is oxygen atom, sulfur atom,  $-N(methyl)-$ , or  $-N(ethyl)-$ ,  $A^1$  is ethylene),  $R^2$  and  $R^3$  independently represent hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, dimethylamino group, acetylamino group, or methylsulfonylamino group (provided that when  $Q$  is phenyl group,  $A^1$  is a single bind or unsubstituted methylene, and  $A^2$  is a single bind, one of  $R^2$  and  $R^3$  is a substituent other than hydrogen atom),  $p$  in  $Rc$  is an integer of 2 or 3,  $A^4$  is a single bind or methylene,  $A^5$  is  $-C(O)-$ ,  $-C(S)-$ , or  $-S(O)_2-$ ,  $Rd$  is hydrogen atom, or methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group, cyclopropylmethyl group, cyclopentyl group, cyclopentylmethyl group, cyclohexyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, or pyridin-4-yl group,  $Re$  is methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, phenylmethyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-

3-yl group, pyridin-4-yl group, furan-2-yl group, furan-3-yl group, thiophen-2-yl group, thiophen-3-yl group, methoxy group, ethoxy group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenyloxy group, 4-methylphenyloxy group, 4-chlorophenyloxy group, 4-fluorophenyloxy group, thiomethoxy group, amino group, N-methylamino group, N,N-dimethylamino group, N-ethylamino group, N,N-diethylamino group, N-propylamino group, N-isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group, N-(4-fluorophenyl)amino group, N-(pyridin-2-yl)amino group, N-(pyridin-3-yl)amino group, N-(pyridin-4-yl)amino group, N-(furan-2-yl)amino group, N-(furan-3-yl)amino group, N-(thiophen-2-yl)amino group, N-(thiophen-3-yl)amino group, pyrrolidino group, piperidino group, morpholino group, methyloxycarbonylamino group or ethyloxycarbonylamino group, R<sub>z</sub>' is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl

group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, 2-(N-ethyl-N-phenylamino)ethyl group, isobutyryl group, isopropylthiocarbonyl group, isopropylsulfonyl group, valeryl group, butylthiocarbonyl group, isovaleryl group, isobutylthiocarbonyl group, pivaloyl group, t-butylthiocarbonyl group, cyclopropylcarbonyl group, cyclopropylthiocarbonyl group, cyclopentylcarbonyl group, cyclopentylthiocarbonyl

group, cyclohexylcarbonyl group, cyclohexylthiocarbonyl group, cyclopentylmethylcarbonyl group, cyclopentylmethylthiocarbonyl group, cyclohexylmethylcarbonyl group, cyclohexylmethylthiocarbonyl group, benzoyl group, thiobenzoyl group, phenylsulfonyl group, 4-methylphenylcarbonyl group, 4-methylphenylthiocarbonyl group, 4-methylphenylsulfonyl group, 4-chlorophenylcarbonyl group, 4-chlorophenylthiocarbonyl group, 4-fluorophenylcarbonyl group, 4-fluorophenylthiocarbonyl group, isopropoxy carbonyl group, N-isopropyl carbamoyl group, N-isopropylthiocarbamoyl group, butyloxy carbonyl group, N-butyl carbamoyl group, N-butylthiocarbamoyl group, isobutyloxy carbonyl group, N-isobutyl carbamoyl group, N-isobutylthiocarbamoyl group, t-butyloxy carbonyl group, N-t-butyl carbamoyl group, N-t-butylthiocarbamoyl group, cyclopropyloxy carbonyl group, N-cyclopropyl carbamoyl group, N-cyclopropylthiocarbamoyl group, cyclopentyloxy carbonyl group, N-cyclopentyl carbamoyl group, N-cyclopentylthiocarbamoyl group, cyclohexyloxy carbonyl group, N-cyclohexyl carbamoyl group, N-cyclohexylthiocarbamoyl group, cyclopentylmethyloxy carbonyl group, cyclohexylmethyloxy carbonyl group, phenyloxy carbonyl group, N-phenyl carbamoyl group, N-phenylthiocarbamoyl group, 4-methylphenyloxy carbonyl group, N-(4-methylphenyl) carbamoyl group, N-(4-methylphenyl)thiocarbamoyl group, 4-chlorophenyloxy carbonyl group, N-(4-chlorophenyl) carbamoyl group, N-(4-chlorophenyl)thiocarbamoyl group, 4-fluorophenyloxy carbonyl group, N-(4-fluorophenyl) carbamoyl group, N-(4-fluorophenyl)thiocarbamoyl group, (pyrrolidino-1-yl) carbonyl group, (piperidino-1-yl) carbonyl group, or (morpholino-4-yl) carbonyl group, Ry' is hydrogen atom, methyl group, ethyl group or isobutyl group, or binds to Rz' to form pyrrolidino group, piperidino group, piperazino group, morpholino group, pyrrol-1-yl group, imidazol-1-yl group, or pyrazol-1-yl group together with the nitrogen atom, provided that

when -D-Rx' or -N(Ry')(Rz') contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when the substituent -D-Rx' or -N(Ry')(Rz') contains amino group, the amino group may be protected with Rp<sup>2</sup>,

AR' is naphthalen-2-yl group, naphthalen-1-yl group, benzofuran-5-yl group, benzofuran-4-yl group, benzofuran-2-yl group, benzo[b]thiophen-5-yl group, benzo[b]thiophen-4-yl group, benzo[b]thiophen-2-yl group, indol-5-yl group, indol-4-yl group, indol-6-yl group, benzothiazol-6-yl group, benzothiazol-7-yl group, benzothiazol-5-yl group, benzothiazol-4-yl group, dihydro-3H-benzothiazol-6-yl group, dihydro-3H-benzothiazol-7-yl group, dihydro-3H-benzothiazol-5-yl group, dihydro-3H-benzothiazol-4-yl group, quinolin-6-yl group, quinolin-3-yl group, quinolin-5-yl group, quinolin-7-yl group, dihydro-1H-quinolin-6-yl group, dihydro-1H-quinolin-5-yl group, benzo[d]isothiazol-5-yl group, benzo[d]isothiazol-4-yl group, benzo[d]isothiazol-6-yl group, benzo[d]isothiazol-7-yl group, 1H-indazol-5-yl group, 1H-indazol-4-yl group, 1H-indazol-6-yl group, benzo[c]isothiazol-5-yl group, benzo[c]isothiazol-4-yl group, benzo[c]isothiazol-6-yl group, benzo[c]isothiazol-7-yl group, 2H-indazol-5-yl group, 2H-indazol-4-yl group, 2H-indazol-6-yl group, imidazo[1,2-a]pyridin-6-yl group, imidazo[1,2-a]pyridin-7-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1H-pyrrolo[2,3-b]pyridin-4-yl group, isoquinolin-6-yl group, isoquinolin-3-yl group, isoquinolin-5-yl group, isoquinolin-7-yl group, dihydro-2H-isoquinolin-6-yl group, dihydro-2H-isoquinolin-5-yl group, cinnolin-6-yl group, cinnolin-5-yl group, quinazolin-6-yl group, quinazolin-7-yl group, quinazolin-5-yl group, quinoxalin-2-yl group, quinoxalin-6-yl group, quinoxalin-5-yl group, 1H-benzimidazol-5-yl group, 1H-benzimidazol-4-yl group, benzoxazol-5-yl group, benzoxazol-6-yl group, benzoxazol-4-yl group, benzoxazol-7-yl group, 1H-pyrrolo[3,2-b]pyridin-5-yl group, 1H-pyrrolo[3,2-b]pyridin-6-yl group, benzo[1,2,5]thiadiazol-5-yl group, benzo[1,2,5]thiadiazol-4-yl group, 1H-benzotriazol-5-yl group, 1H-benzotriazol-4-yl group, 1,3-dihydropyrrolo[2,3-

b]pyridin-5-yl group, 1,3-dihydropyrrolo[2,3-b]pyridin-4-yl group, 1,3-dihydrobenzimidazol-5-yl group, 1,3-dihydrobenzimidazol-4-yl group, dihydro-3H-benzoxazol-6-yl group, dihydro-3H-benzoxazol-7-yl group, dihydro-3H-benzoxazol-5-yl group, dihydro-3H-benzoxazol-4-yl group, phthalazin-6-yl group, phthalazin-5-yl group, [1,8]naphthalidin-3-yl group, [1,8]naphthalidin-4-yl group, [1,5]naphthalidin-3-yl group, [1,5]naphthalidin-4-yl group, 1H-pyrrolo[3,2-c]pyridin-6-yl group, 1H-pyrrolo[3,2-c]pyridin-4-yl group, 1H-pyrrolo[2,3-c]pyridin-5-yl group, 1H-pyrrolo[2,3-c]pyridin-4-yl group, 1H-pyrazolo[4,3-b]pyridin-5-yl group, 1H-pyrazolo[4,3-b]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-4-yl group, 1H-pyrazolo[3,4-c]pyridin-5-yl group, 1H-pyrazolo[3,4-c]pyridin-4-yl group, 1H-pyrazolo[3,4-b]pyridin-5-yl group, 1H-pyrazolo[3,4-b]pyridin-4-yl group, [1,2,4]triazolo[4,3-a]pyridin-6-yl group, [1,2,4]triazolo[4,3-a]pyridin-7-yl group, thieno[3,2-c]pyridin-2-yl group, thieno[3,2-c]pyridin-3-yl group, thieno[3,2-c]pyridin-6-yl group, thieno[3,2-b]pyridin-2-yl group, thieno[3,2-b]pyridin-3-yl group, thieno[3,2-b]pyridin-5-yl group, thieno[3,2-b]pyridin-6-yl group, 1H-thieno[3,2-c]pyrazol-5-yl group, 1H-thieno[3,2-c]pyrazol-4-yl group, benzo[d]isoxazol-5-yl group, benzo[d]isoxazol-4-yl group, benzo[d]isoxazol-6-yl group, benzo[d]isoxazol-7-yl group, benzo[c]isoxazol-5-yl group, benzo[c]isoxazol-4-yl group, benzo[c]isoxazol-6-yl group, benzo[c]isoxazol-7-yl group, indolizin-7-yl group, indolizin-6-yl group, indolizine-8-yl group, 1,3-dihydroindol-5-yl group, 1,3-dihydroindol-4-yl group, 1,3-dihydroindol-6-yl group, 1H-pyrazolo[3,4-d]thiazol-5-yl group, 2H-isoindol-5-yl group, 2H-isoindol-4-yl group, [1,2,4]triazolo[1,5-a]pyrimidin-6-yl group, 1H-pyrazolo[3,4-b]pyrazin-5-yl group, 1H-imidazo[4,5-b]pyrazin-5-yl group, 7H-purin-2-yl group, 4H-chromen-6-yl group, or 4H-chromen-5-yl group (the aforementioned groups may be substituted with one of Xa or two or more of the same or different Xa), and Xa is oxo group, thioxo group, fluorine atom, chlorine atom, trifluoromethyl group, methyl group, ethyl group,

propyl group, 2-hydroxyethyl group, carboxymethyl group, 2-carboxyethyl group, N,N-dimethylcarbamoylmethyl group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, carboxymethyloxy group, 2-carboxyethyloxy group, N,N-dimethylcarbamoylmethyloxy group, amino group, methylamino group, dimethylamino group, 2-hydroxyethylamino group, carbamoylamino group, acetylamino group, furan-2-carboxyamino group, 2-hydroxyacetylamino group, 2-aminoacetylamino group, methylsulfonylamino group, (N,N-dimethylsulfamoyl)amino group, methanesulfonyl group, sulfamoyl group, N-methylsulfamoyl group, N,N-dimethylsulfamoyl group, carboxyl group, acetyl group, carbamoyl group, or N,N-dimethylcarbamoyl group, provided that when AR' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when AR' contains amino group, the amino group may be protected with Rp<sup>2</sup>.

(179) The compound according to (119) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5'</sup> is a ring-constituting carbon atom substituted with Zx', or an unsubstituted ring-constituting carbon atom, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms, and Rs' is -O-Rx'.

(180) The compound according to (131) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5'</sup> is a ring-constituting carbon atom substituted with Zx, or an unsubstituted ring-constituting carbon atom, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms, and D is oxygen atom.

(181) The compound according to (132) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5'</sup> is a ring-constituting carbon atom substituted with Zx', or an unsubstituted ring-constituting carbon atom, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms, and Rs' is -O-Rx'.



(182) The compound according to (129) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5'</sup> is carbon atom substituted with nitro group, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms, and Rs' is -O-Rx'.

(183) The compound according to (131) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5'</sup> is carbon atom substituted with nitro group, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms, and Rs' is -O-Rx'.

(184) The compound according to (132) mentioned above, wherein, in the formula (III), AR' binds to C<sup>3'</sup> in the aromatic ring (E'), Rs' binds to C<sup>4'</sup> in the aromatic ring (E'), C<sup>5'</sup> is carbon atom substituted with nitro group, C<sup>2'</sup> and C<sup>6'</sup> are unsubstituted ring-constituting carbon atoms, and Rs' is -O-Rx'.

(185) An agent for prophylactic and/or therapeutic treatment of fibrosis, which contains a type 4 PLA<sub>2</sub> inhibitor as an active ingredient.

(186) An agent for prophylactic and/or therapeutic treatment of pulmonary fibrosis, which contains a type 4 PLA<sub>2</sub> inhibitor as an active ingredient..

(187) The prophylactic and/or therapeutic agent according to (186), wherein the pulmonary fibrosis is drug-induced pulmonary fibrosis.

(188) The prophylactic and/or therapeutic agent according to (187), wherein the drug-induced pulmonary fibrosis is a disease induced by one or more kinds of medicaments among methotrexate, sodium aurothiomalate, auranofin, D-penicillamine, bucillamine, actarit, salazosulfapyridine, cyclophosphamide, Taxol, etoposide, cisplatin, vincristine, vinblastine, irinotecan, gefitinib, and bleomycin.

(189) The prophylactic and/or therapeutic agent according to (187), wherein the drug-induced pulmonary fibrosis is a disease induced by one or more kinds of medicaments among methotrexate and bleomycin.

(190) The prophylactic and/or therapeutic agent according to (186), wherein the

type 4 PLA<sub>2</sub> inhibitor is a compound represented by the formula (I) or a pharmacologically acceptable salt thereof.

(191) The prophylactic and/or therapeutic agent according to (186), wherein the type 4 PLA<sub>2</sub> inhibitor is an inhibitor selected from the group consisting of 4-(1-benzhydryl-6-chloro-1H-indol-3-ylmethyl)-3-methoxybenzoic acid, 4-{4-[2-(2-[bis(4-chlorophenyl)methoxy]ethylsulfonyl)ethoxy]phenyl}-1,1,1-trifluoro-2-butanone, N-{1-[2-(2,4-difluorobenzoyl)benzoyl]-4-tritylsulfanylpyrrolidin-2-ylmethyl}-4-(2,4-dioxothiazolidin-5-ylidenemethyl)benzoic acid amide, 4-methyl-2-oxo-5-(5,6,7,8-tetrahydronaphthalen-2-yl)oxazolidine-3-carboxylic acid (6-methoxytetrahydropyran-2-yl)amide, 4-methyl-2-oxo-5-(4-methylphenyl)thiazolidine-3-carboxylic acid (tetrahydropyran-2-yl)amide, 4-[3-(4-decyloxyphenoxy)-2-oxopropoxy]benzoic acid, and 1-{2-[4-(carboxymethyl)phenoxy]ethyl}-3-dodecanoylindole-2-carboxylic acid.

The compound (I) of the present invention or a pharmaceutically acceptable salt thereof has an action of suppressing the production of both of prostaglandins and leukotrienes, and said compound has characteristic features that, when administered to a human or animal, the compound exerts superior prophylactic and/or therapeutic effect on diseases or pathological conditions in which a prostaglandin and/or leukotriene is involved, and the compound has extremely low toxicity. The compounds (II) and (III) of the present invention are synthetic intermediates useful for the production of the compound (I) of the present invention. Furthermore, it was confirmed that a type 4 PLA<sub>2</sub> inhibitor is useful as a prophylactic and/or therapeutic agent for fibrosis, in particular, pulmonary fibrosis, especially drug-induced pulmonary fibrosis, which was induced as a side effect of a medicament.

## Best Mode for Carrying out the Invention

In the present specification, carbon atom may sometimes be represented simply by "C", hydrogen atom by "H", oxygen atom by "O", sulfur atom by "S", and nitrogen atom by "N".

Examples of Link in the aforementioned general formula (I) include a saturated straight hydrocarbon chain having 1 to 3 carbon atoms or an unsaturated straight hydrocarbon chain having 2 or 3 carbon atoms. In the present invention, the straight chain of the saturated straight hydrocarbon chain is preferably unsubstituted. The straight chain of the unsaturated straight hydrocarbon chain is also preferably unsubstituted. As the saturated straight hydrocarbon chain,  $-(CH_2)_n-$  is preferred. Symbol  $n$  is an integer of 1 to 3. When  $n$  is 1, 2 or 3, the desired action is most characteristically exhibited. Methylene where  $n$  is 1, ethylene where  $n$  is 2 and trimethylene where  $n$  is 3 are preferred, and ethylene where  $n$  is 2 is particularly preferred.

The unsaturated hydrocarbon chain having 2 or 3 carbon atoms means a hydrocarbon chain which contains an unsaturated bond as a double bond or a triple bond among the carbon-carbon bonds. As the unsaturated hydrocarbon chain, an unsaturated hydrocarbon chain containing a double bond is preferred. When the chain contains one or more double bonds, the number of the double bond may preferably one. Specific examples include ethenylene which has two carbon atoms and contains one double bond, as well as ethynylene which has two carbon atoms and contains one triple bond, propen-3-yl which has three carbon atoms and contains one double bond, and propyn-3-yl which has three carbon atoms and contains one triple bond.

$C^2$ ,  $C^3$ ,  $C^4$ ,  $C^5$  and  $C^6$  in the aromatic ring (E) in the formula (I) each represent a ring-constituting carbon atom. The ring-constituting carbon atoms form the aromatic ring (E), and accordingly, they are represented as C or CH.

Among them, any one of ring-constituting carbon atoms to which Rs or Ar does not bind may be replaced with V. The aforementioned expression "to be replaced with" means that any one of the ring-constituting carbon atoms C<sup>2</sup>, C<sup>3</sup>, C<sup>4</sup>, C<sup>5</sup> and C<sup>6</sup> is replaced with V, and thus V may sometimes be a ring-constituting component. Rs and AR each bind to any of the ring-constituting carbon atoms C<sup>2</sup>, C<sup>3</sup>, C<sup>4</sup>, C<sup>5</sup> or C<sup>6</sup> in the aromatic ring (E), and this means that, for example, when AR binds to C<sup>2</sup>, Rs binds to any of the ring-constituting carbon atoms C<sup>3</sup>, C<sup>4</sup>, C<sup>5</sup> and C<sup>6</sup>, when AR binds to C<sup>3</sup>, Rs binds to any of the ring-constituting carbon atoms C<sup>2</sup>, C<sup>4</sup>, C<sup>5</sup> and C<sup>6</sup>, and when AR bind to C<sup>4</sup>, Rs binds to the ring-constituting carbon atom C<sup>2</sup> or C<sup>3</sup>. Preferred examples of these combinations of substitution positions include a compound wherein AR binds to C<sup>2</sup>, and Rs binds to any of the atoms C<sup>3</sup>, C<sup>4</sup>, and C<sup>5</sup>, and particularly preferred examples include a compound wherein AR binds to C<sup>2</sup>, and Rs binds to C<sup>3</sup> or C<sup>4</sup>. Preferred examples also include a compound wherein AR binds to C<sup>3</sup>, and Rs binds to any of the atoms C<sup>4</sup>, C<sup>5</sup>, and C<sup>6</sup>, and particularly preferred examples also include a compound wherein AR binds to C<sup>3</sup>, and Rs binds to the atom C<sup>4</sup> or C<sup>5</sup>. A still more preferred example is a compound wherein AR binds to C<sup>3</sup>, and Rs binds to C<sup>4</sup>.

One of the atoms C<sup>2</sup>, C<sup>3</sup>, C<sup>4</sup>, C<sup>5</sup> and C<sup>6</sup> to which Rs and AR do not bind may be replaced with V. For example, when AR binds to C<sup>2</sup>, and Rs binds to C<sup>3</sup>, one of the ring-constituting carbon atoms C<sup>4</sup>, C<sup>5</sup>, and C<sup>6</sup> may be replaced with V. As another example, it is meant that when AR binds to C<sup>3</sup>, and Rs binds to C<sup>4</sup>, one of the atoms C<sup>2</sup>, C<sup>5</sup>, and C<sup>6</sup> may be replaced with V. Among them combinations and other combinations, preferred examples are a compound wherein AR binds to C<sup>2</sup>, Rs binds to C<sup>3</sup>, and C<sup>4</sup> is replaced with V; a compound wherein AR binds to C<sup>2</sup>, Rs binds to C<sup>4</sup>, and C<sup>5</sup> is replaced with V; a compound wherein AR binds to C<sup>2</sup>, Rs binds to C<sup>5</sup>, and C<sup>4</sup> is replaced with V; a compound wherein AR binds to C<sup>3</sup>, Rs binds to C<sup>4</sup>, and C<sup>5</sup> is replaced with V; a compound wherein AR binds to C<sup>3</sup>, Rs binds to C<sup>4</sup>, and C<sup>6</sup> is

replaced with V; a compound wherein AR binds to C<sup>3</sup>, Rs binds to C<sup>5</sup>, and C<sup>4</sup> is replaced with V; a compound wherein AR binds to C<sup>3</sup>, Rs binds to C<sup>6</sup>, and C<sup>5</sup> is replaced with V, and the like. Furthermore, particularly preferred examples include a compound wherein AR binds to C<sup>3</sup>, Rs binds to C<sup>4</sup>, and C<sup>5</sup> is replaced with V; and a compound wherein AR binds to C<sup>3</sup>, Rs binds to C<sup>4</sup>, and C<sup>6</sup> is replaced with V, and an particularly preferred example is a compound wherein AR binds to C<sup>3</sup>, Rs binds to C<sup>4</sup>, and C<sup>5</sup> is replaced with V.

V represents nitrogen atom, or carbon atom substituted with Zx. Namely, when V represent nitrogen atom, the aromatic ring (E) in the formula (I) represents a pyridine ring. When V represent carbon atom substituted with Zx, the aromatic ring (E) is a benzene ring having Zx. Both of the compounds are particularly preferred. Furthermore, a compound wherein AR binds to C<sup>3</sup>, Rs binds to C<sup>4</sup>, C<sup>5</sup> is V replaced with V, and this V represents nitrogen atom is particularly preferred.

Zx is defined as a linear or branched saturated alkyl group having 1 to 4 carbon atoms, fluorine atom, chlorine atom, bromine atom, nitro group, -OR<sup>9</sup>, or -N(R<sup>n1</sup>)(R<sup>n2</sup>). Among them, fluorine atom, chlorine atom, bromine atom, and nitro group are preferred examples, and fluorine atom is particularly preferred.

As for Zx, examples of the linear or branched saturated alkyl group having 1 to 4 carbon atoms include methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group and the like, and among them, methyl group is particularly preferred.

R<sup>9</sup> represents hydrogen atom, a lower alkyl group having 1 to 4 carbon atoms, or -A<sup>6</sup>-Qp. Among them, hydrogen atom is a particularly preferred example. Preferred examples of the lower alkyl group having 1 to 4 carbon atoms include methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, and the like, and methyl group is particularly preferred.

A<sup>6</sup> in -A<sup>6</sup>-Qp represents a single bond or methylene, and Qp represents a

phenyl group which may be substituted with one of  $T^1$  or two or more of the same or different  $T^1$ . The substituent  $T^1$  is a linear or branched saturated alkyl group having 1 to 4 carbon atoms, hydroxyl group, fluorine atom, chlorine atom, bromine atom, trifluoromethyl group, nitro group, an alkoxy group having 1 to 4 carbon atoms, or a mono- or dialkylamino group having 1 to 4 carbon atoms. Specific examples of  $-A^6-Qp$  include phenyl group, methylphenyl group, chlorophenyl group, benzyl group, methylbenzyl group, chlorobenzyl group, dichlorobenzyl group, fluorobenzyl group, trifluoromethylbenzyl group, nitrobenzyl group, methoxyphenyl group, N-methylaminobenzyl group, N,N-dimethylaminobenzyl group, and the like.

Preferred examples of  $-OR^9$  include hydroxyl group, methoxy group, and the like, and hydroxyl group is particularly preferred.

$Rn^1$  represents hydrogen atom or a linear or branched saturated alkyl group having 1 to 4 carbon atoms, and hydrogen atom is particularly preferred. Examples of the linear or branched saturated alkyl group having 1 to 4 carbon atoms include methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, or t-butyl group, and the like. Among them, methyl group, ethyl group, propyl group, isopropyl group, and the like are preferred examples, and methyl group is particularly preferred.

$Rn^2$  has the same meaning as  $Rn^1$ , or represents a  $-COR^{23}$  group or a  $-SO_2R^{24}$  group, or binds to  $Rn^1$  to form a 3- to 6-membered ring together with the nitrogen atom to which they bind to form a saturated nitrogen-containing cycloalkyl group or morpholino group.

$R^{23}$  represents hydrogen atom, a lower alkyl group having 1 to 4 carbon atoms, a lower alkoxy group having 1 to 4 carbon atoms,  $-O-A^6-Qp$ , or  $-N(R^{25})(R^{26})$ .  $R^{25}$  represents hydrogen atom, or a linear or branched saturated alkyl group having 1 to 4 carbon atoms.  $R^{26}$  has the same meaning as  $R^{25}$ , or binds to  $R^{25}$  to form a 3- to 6-membered ring together with the nitrogen atom to which they bind to represent a

saturated nitrogen-containing cycloalkyl group or morpholino group. Examples of the compound wherein  $R^{26}$  "binds to  $R^{25}$  to form a 3- to 6-membered ring together with the nitrogen atom to which they bind to represent a saturated nitrogen-containing cycloalkyl group or morpholino group" include, for example, a compound wherein a cyclic aminoalkyl group containing nitrogen atom such as pyrrolidino group, piperazino group and morpholino group is formed.

Specific examples of  $-COR^{23}$  include formyl group, acetyl group, t-butyloxycarbonyl group, phenyloxycarbonyl group, benzyloxycarbonyl group, carbamoyl group, N-methylcarbamoyl group, N,N-dimethylcarbamoyl group, piperidine-1-carbonyl group, morpholine-4-carbonyl group, and the like, and preferred examples include formyl group, acetyl group, carbamoyl group, and the like. In the aforementioned formulas, as represented by  $A^6$  and  $Qp$ , for example, the same symbols may sometimes be used simultaneously at different positions. These symbols are used to mean the same class of groups of substituents. However, because each substituent is independently chosen from each other, the same symbols do not mean that an identical substituent should be necessarily chosen, and as a result, selection of the same or different kind of substituent is not prohibited.

$R^{24}$  represents a lower alkyl group having 1 to 4 carbon atoms, amino group, or a mono- or dialkylamino group having 1 to 4 carbon atoms. Specific examples of  $-SO_2R^{24}$  include mesyl group, sulfamoyl group, N-methylsulfamoyl group, N,N-dimethylsulfamoyl group, and the like, and preferred examples include mesyl group, N,N-dimethylsulfamoyl group, and the like.

Specific examples of  $-N(R^{n1})(R^{n2})$  include amino group, N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, piperidino group, pyrrolidino group, morpholino group, formylamino group, acetylamino group, t-butyloxycarbonylamino group, phenyloxycarbonylamino group,

benzyloxycarbonylamino group, carbamoylamino group, N-methylcarbamoylamino group, N,N-dimethylcarbamoylamino group, piperidine-1-carbonylamino group, morpholine-4-carbonylamino group, mesylamino group, sulfamoylamino group, N-methylsulfamoylamino group, N,N-dimethylsulfamoylamino group, and the like. Among them, preferred examples include amino group, N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, N,N-dimethylsulfamoylamino group, and the like, and amino group, N-methylamino group, and N,N-dimethylamino group are particularly preferred.

Therefore, preferred examples of Zx include fluorine atom, chlorine atom, bromine atom, nitro group, methyl group, hydroxyl group, methoxy group, amino group, N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, N,N-dimethylsulfamoylamino group, and the like, and particularly preferred examples include fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, N,N-dimethylamino group, and the like.

In the formula (I), Rs is defined to represent -D-Rx or -N(Ry)(Rz).

D is defined to represent a single bond, oxygen atom, sulfur atom, -S(O)-, -S(O)<sub>2</sub>-, or -C(O)-. Among them, oxygen atom and sulfur atom are preferred, and oxygen atom is particularly preferred. Another preferred examples include the compounds wherein D represent a single bond.

Rx represents a linear or branched saturated alkyl group having 3 to 8 carbon atoms, or represents Ra, Rb, or Rc mentioned above.

As for Rx, examples of the linear or branched saturated alkyl group having 3 to 8 carbon atoms include, for example, propyl group, isopropyl group, butyl group,



isobutyl group, 1-methylpropyl group, t-butyl group, pentyl group, isopentyl group, 2-methylbutyl group, 2,2-dimethylpropyl group, hexyl group, 4-methylpentyl group, 2,3-dimethylbutyl group, 2-ethylbutyl group, heptyl group, octyl group, and the like, and butyl group, isobutyl group, and 2-ethylbutyl group are particularly preferred.

As for  $R_x$ ,  $R^1$  of  $R_a$  is defined to be a saturated cyclic alkyl group having 3 to 7 carbon atoms substituted with a lower alkyl group having 1 to 4 carbon atoms or an unsubstituted saturated cyclic alkyl group having 3 to 7 carbon atoms, or a condensed saturated cyclic alkyl group having 6 to 8 carbon atoms substituted with a lower alkyl group having 1 to 4 carbon atoms or an unsubstituted condensed saturated cyclic alkyl group having 6 to 8 carbon atoms. As for  $R^1$ , examples of the saturated cyclic alkyl group having 3 to 7 carbon atoms include cyclopropyl group, cyclobutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, and the like, and cyclopentyl group, cyclohexyl group, and cycloheptyl group are particularly preferred. As for  $R^1$ , examples of the condensed saturated cyclic alkyl group having 6 to 8 carbon atoms group include bicyclo[2,2,1]heptyl group, bicyclo[2,2,2]octyl group, and the like.

Examples of the lower alkyl group having 1 to 4 carbon atoms substituting on  $R^1$  include methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, and the like. Examples of  $R^1$  substituted with a lower alkyl group having 1 to 4 carbon atoms include methylcyclopentyl group, methylcyclohexyl group, methylbicyclo[2,2,1]heptyl group, and the like.

Symbol  $k$  is defined to be 0 or an integer of 1 to 3. A single bond where  $k$  is 0, methylene where  $k$  is 1, and ethylene where  $k$  is 2 are preferred, and a bond where  $k$  is 0, and methylene where  $k$  is 1 are particularly preferred.

Examples of  $R_a$  include cyclopropyl group, cyclobutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopropylmethyl group, cyclobutylmethyl group, cyclopentylmethyl group, cyclohexylmethyl group,

cycloheptylmethyl group, 2-cyclopentylethyl group, 2-cyclohexylethyl group, 3-cyclohexylpropyl group, 2-methylcyclopentyl group, 3-methylcyclopentyl group, 3,4-dimethylcyclopentyl group, 4-methylcyclohexyl group, 4,4-dimethylcyclohexyl group, 4-ethylcyclohexyl group, 4-methylcyclohexylmethyl group, bicyclo[2,2,1]heptane-2-methyl group, bicyclo[2,2,2]octane-2-methyl group, 3-methylbicyclo[2,2,1]heptane-2-methyl group, bicyclo[2,2,1]hept-1-ylmethyl group, bicyclo[2,2,2]oct-1-ylmethyl group, and the like. Cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-cyclopentylethyl group, 2-cyclohexylethyl group are preferred, and cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group are particularly preferred.

As for Rx, A<sup>2</sup> in Rb is defined to be a single bond, oxygen atom, sulfur atom, -S(O)-, -S(O)<sub>2</sub>-, or -N(R<sup>4</sup>)-. R<sup>4</sup> is defined to be a lower alkyl group having 1 to 4 carbon atoms. Preferred examples are methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, and the like, and methyl group and ethyl group are particularly preferred examples. Therefore, particularly preferred examples of A<sup>2</sup> include a single bond, oxygen atom, sulfur atom, -N(methyl)-, and -N(ethyl)-.

A<sup>1</sup> is defined to be a single bond or an alkylene (a) having 1 to 3 carbon atoms, i.e., methylene, ethylene or trimethylene. However, when A<sup>2</sup> represents oxygen atom, sulfur atom, -S(O)-, -S(O)<sub>2</sub>- or -N(R<sup>4</sup>)-, A<sup>1</sup> is ethylene or trimethylene. Further, the alkylene (a) may be substituted with a lower alkyl group having 1 to 4 carbon atoms or phenyl group. Examples of the lower alkyl group having 1 to 4 carbon atoms for the above compound include methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, and the like, and methyl group, and ethyl group are preferred examples. Specific examples of A<sup>1</sup> include methylene, methylmethylene, ethylmethylene, phenylmethylene, ethylene,

methylethylene, dimethylethylene, ethylethylene, phenylethylene, trimethylene, methyltrimethylene, and the like. Among them, when A<sup>2</sup> represents a single bond, A<sup>1</sup> is most preferably a single bond, or methylene, methylmethylene, or ethylene. Further, when A<sup>2</sup> represents oxygen atom, sulfur atom, -S(O)-, -S(O)<sub>2</sub>- or -N(R<sup>4</sup>)-, A<sup>1</sup> is most preferably ethylene.

Q in Rb is defined to be a residue of a partially unsaturated or completely unsaturated monocyclic or condensed bicyclic carbon ring or heterocyclic ring (q), and the heterocyclic ring (q) means a ring containing 1 to 4 the same or different ring-constituting heteroatoms selected from the group consisting of nitrogen atom, oxygen atom, and sulfur atom. The term "residue" means a monovalent group formed by eliminating hydrogen atom bonding to a ring-constituting atom. The residue of monocyclic carbon ring or heterocyclic ring is a partially unsaturated or completely unsaturated substituent having 5 to 7 atoms, and examples include, for example, phenyl group, thienyl group, furyl group, pyrrolyl group, pyridyl group, oxazolyl group, isoxazolyl group, thiazolyl group, isothiazolyl group, imidazolyl group, pyrazolyl group, oxadiazolyl group, thiadiazolyl group, triazolyl group, tetrazolyl group, and the like. Among them, phenyl group, thienyl group, furyl group, pyridyl group, and oxazolyl group are preferred examples, and phenyl group is particularly preferred.

The condensed bicyclic carbon ring or heterocyclic ring is a partially unsaturated or completely unsaturated ring having 8 to 11 atoms, and examples of residue thereof include, for example, naphthyl group, tetrahydronaphthyl group, indanyl group, indenyl group, quinolyl group, isoquinolyl group, indolyl group, benzofuryl group, benzothienyl group, benzimidazolyl group, benzoxazolyl group, benzothiazolyl group, indazolyl group, 4H-chromenyl group, dihydrobenzodioxyl group, benzoisoxazolyl group, pyrrolopyridinyl group, pyrazolopyridinyl group, triazolopyridinyl group, thienopyridinyl group, thienopyrazolyl group,

1,3-dihydrobenzimidazole group, dihydro-3H-benzoxazole group, dihydro-3H-benzothiazole group, and the like. Among them, naphthyl group, tetrahydronaphthyl group, indanyl group, indolyl group, and dihydrobenzodioxyl group are preferred examples, and indanyl group is one of particularly preferred examples.

Q binds to A<sup>2</sup> at an arbitrary position on the ring. Preferred examples of Q with indication of bonding position include phenyl group, 2- or 3-thienyl group, 2- or 3-furyl group, 2-, 3- or 4-pyridyl group, 2-, 4- or 5-oxazolyl group, 1- or 2-naphthyl group, 1-, 2-, 5- or 6-tetrahydronaphthyl group, indan-1-yl group, indan-2-yl group, indan-4-yl group, indan-5-yl group, 1-, 2-, 3-, 4-, 5-, 6-, or 7-indolyl group, 2-, 5- or 6-dihydrobenzodioxyl group, and the like. Among them, phenyl group, and indan-2-yl group are particularly preferred.

In R<sup>b</sup>, R<sup>2</sup> and R<sup>3</sup> are defined to be substituents of Q, and independently represent hydrogen atom, a linear or branched saturated alkyl group having 1 to 4 carbon atoms, oxo group, thioxo group, fluorine atom, chlorine atom, bromine atom, trifluoromethyl group, -OR<sup>5</sup>, -N(R<sup>6</sup>)(R<sup>6'</sup>), -NHCOR<sup>7</sup>, -NHSO<sub>2</sub>R<sup>8</sup>, or -A<sup>6</sup>-Qa, or bind to each other to represent methylenedioxy group.

Examples of the linear or branched saturated alkyl group having 1 to 4 carbon atoms include methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, and the like, and methyl group is particularly preferred.

R<sup>6</sup> in -N(R<sup>6</sup>)(R<sup>6'</sup>) represents hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms. R<sup>6'</sup> has the same meaning as R<sup>6</sup>, or binds to R<sup>6</sup> to form a 3- to 6-membered ring together with the nitrogen atom to which they bind to form a saturated nitrogen-containing cycloalkyl group or morpholino group. Therefore, specific examples of -N(R<sup>6</sup>)(R<sup>6'</sup>) include amino group, N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group,

N,N-dimethylamino group, N,N-diethylamino group, piperidino group, pyrrolidino group, morpholino group, and the like. N,N-Dimethylamino group, piperidino group, morpholino group, and the like are preferred examples, and N,N-dimethylamino group is a particularly preferred example.

R<sup>5</sup> and R<sup>7</sup> are defined to independently represent hydrogen atom, a lower alkyl group having 1 to 4 carbon atoms, or a -A<sup>6</sup>-Qa group. Examples of the lower alkyl group having 1 to 4 carbon atoms include methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, and the like, and among them, methyl group is a preferred example.

A<sup>6</sup> in -A<sup>6</sup>-Qa has the same meaning as that defined above. Qa is defined to be a partially unsaturated or completely unsaturated monocyclic or condensed bicyclic carbon ring or heterocyclic ring (qa), and the heterocyclic ring (qa) means a substituent containing 1 to 4 the same or different ring-constituting heteroatoms selected from the group consisting of nitrogen atom, oxygen atom, and sulfur atom. The monocyclic carbon ring or heterocyclic ring is a partially unsaturated or completely unsaturated ring having 5 to 7 atoms, and examples of residue thereof include, for example, phenyl group, thienyl group, furyl group, pyrrolyl group, pyridyl group, oxazolyl group, isoxazolyl group, thiazolyl group, isothiazolyl group, imidazolyl group, pyrazolyl group, oxadiazolyl group, thiadiazolyl group, triazolyl group, tetrazolyl group, and the like. The condensed bicyclic carbon ring or heterocyclic ring is a partially unsaturated or completely unsaturated ring having 8 to 11 atoms, and examples of residue thereof include, for example, naphthyl group, indanyl group, indenyl group, quinolyl group, isoquinolyl group, indolyl group, benzofuryl group, benzothienyl group, benzimidazolyl group, benzoxazolyl group, benzothiazolyl group, indazolyl group, and the like.

Qa binds to A<sup>6</sup> at an arbitrary position on the ring. Further, Qa may be substituted with two or more of the same or different T<sup>1</sup>. T<sup>1</sup> has the same meaning

as defined above.

Specific examples of  $A^6$ -Qa include phenyl group, methylphenyl group, chlorophenyl group, benzyl group, methylbenzyl group, chlorobenzyl group, dichlorobenzyl group, fluorobenzyl group, trifluoromethylbenzyl group, nitrobenzyl group, methoxyphenyl group, N-methylaminobenzyl group, N,N-dimethylaminobenzyl group, furyl group, thienyl group, pyrrolyl group, pyridyl group, oxazolyl group, isoxazolyl group, thiazolyl group, isothiazolyl group, imidazolyl group, pyrazolyl group, oxadiazolyl group, thiadiazolyl group, triazolyl group, tetrazolyl group, naphthyl group, indanyl group, indenyl group, quinolyl group, isoquinolyl group, indolyl group, benzofuryl group, benzothienyl group, benzimidazolyl group, benzoxazolyl group, benzothiazolyl group, indazolyl group, and the like.

$R^8$  each defined to be a lower alkyl group having 1 to 4 carbon atoms, and examples of the lower alkyl group having 1 to 4 carbon atoms include methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, and the like.

Therefore, preferred examples of  $R^2$  and  $R^3$  include hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, dimethylamino group, acetylamino group, and methylsulfonylamino group, and hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, and dimethylamino group are particularly preferred. When Q represents phenyl group,  $A^1$  represents a single bond, or unsubstituted methylene, and  $A^2$  represents a single bond, at least one of  $R^2$  and  $R^3$  preferably represents a substituent other than hydrogen atom.

Particularly preferred examples of Rb include 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl

group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl  
 group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl  
 group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl  
 group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group,  
 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl  
 group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group,  
 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group,  
 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group,  
 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl  
 group, 3-methylphenylmethyl group, 4-methylphenylmethyl group,  
 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group,  
 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl  
 group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group,  
 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group,  
 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group,  
 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group,  
 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group,  
 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group,  
 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group,  
 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group,  
 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group,  
 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group,  
 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group,  
 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group,  
 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,  
 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,  
 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group,

2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, 2-(N-ethyl-N-phenylamino)ethyl group, and the like.

Symbol p in Rc is defined to be an integer of 2 to 4. Ethylene where p is 2, and trimethylene where p is 3 are preferred, and ethylene where p is 2 is particularly preferred. A<sup>4</sup> represents a single bond, or represents methylene or ethylene, and a single bond and methylene are particularly preferred. A<sup>5</sup> represents -C(O)-, -C(S)-, or -S(O)<sub>2</sub>-, and all of them are preferred. Rd represents hydrogen atom, an alkyl group having 1 to 8 carbon atoms, or group Qa. Re represents an alkyl group having 1 to 8 carbon atoms, a -A<sup>6</sup>-Qa group, a -(CH<sub>2</sub>)<sub>i</sub>R<sup>14</sup> group, a -OR<sup>28</sup> group, a -SR<sup>28</sup> group, or a -N(R<sup>29</sup>)(R<sup>30</sup>) group. The group Qa and -A<sup>6</sup>-Qa have the same meanings as defined above.

The alkyl group having 1 to 8 carbon atoms is a linear or branched saturated alkyl group or a linear or branched partially unsaturated alkyl group, or an alkyl group which may contain a cycloalkyl group having 3 to 7 carbon atoms, and examples include, for example, methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, pentyl group, isopentyl group, 2-methylbutyl group, 2,2-dimethylpropyl group, hexyl group, 4-methylpentyl group, 2,3-dimethylbutyl group, 2-ethylbutyl group, heptyl group, octyl group, cyclopropyl group, cyclobutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopropylmethyl group, cyclobutylmethyl group, cyclopentylmethyl group, cyclohexylmethyl group, cycloheptylmethyl group, 2-cyclopentylethyl group, 2-cyclohexylethyl group, 2-methylcyclopentyl group, 3-methylcyclopentyl group, 3,4-dimethylcyclopentyl group, 4-methylcyclohexyl group, 4,4-dimethylcyclohexyl



group, 4-ethylcyclohexyl group, 4-methylcyclohexylmethyl group, and the like.

Symbol  $i$  in  $-(CH_2)_iR^{14}$  represents an integer of 1 to 3, and  $R^{14}$  represents hydroxyl group, an alkoxy group having 1 to 4 carbon atoms, carboxyl group, or an N,N-dialkylcarbamoyl group having 1 to 4 carbon atoms. Examples of the alkoxy group having 1 to 4 carbon atoms include methoxy group, ethoxy group, propyloxy group, isopropyloxy group, butoxy group, isobutyloxy group, t-butyloxy group, and the like. Examples of the N,N-dialkylcarbamoyl group having 1 to 4 carbon atoms include N,N-dimethylcarbamoyl group, N,N-diethylcarbamoyl group, and the like.

$R^{28}$  in  $-OR^{28}$  or  $-SR^{28}$  represents an alkyl group having 1 to 8 carbon atoms, or  $-A^6-Qa$ , and these have the same meanings as defined above.

$R^{29}$  in  $-N(R^{29})(R^{30})$  represents an alkyl group having 1 to 8 carbon atoms, an alkoxycarbonyl group having 1 to 4 carbon atoms, or  $-A^6-Qa$ . Among them, the alkyl group having 1 to 8 carbon atoms and  $-A^6-Qa$  have the same meanings as those defined above. Examples of the alkoxycarbonyl group having 1 to 4 carbon atoms include methyloxycarbonyl group, ethyloxycarbonyl group, propyloxycarbonyl group, isopropyloxycarbonyl group, butyloxycarbonyl group, isobutyloxycarbonyl group, t-butyloxycarbonyl group, and the like.  $R^{30}$  represents hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms, or binds to  $R^{29}$  to form a 3- to 6-membered ring together with the nitrogen atom to which they bind to form a saturated nitrogen-containing cycloalkyl group or morpholino group. The lower alkyl group having 1 to 4 carbon atoms has the same meaning as defined above. Examples of the compound where " $R^{30}$  binds to  $R^{29}$  to form a 3- to 6-membered ring together with the nitrogen atom to which they bind to form a saturated nitrogen-containing cycloalkyl group or morpholino group" include, for example, a compound wherein a cyclic aminoalkyl group containing nitrogen atom such as pyrrolidino group, piperazino group, and morpholino group is formed.

Preferred examples of  $R_d$  include hydrogen atom as well as methyl group,

ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group, cyclopropylmethyl group, cyclopentyl group, cyclopentylmethyl group, cyclohexyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorobenzyl group, 4-fluorobenzyl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, and the like.

Particularly preferred examples of R<sub>d</sub> include methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, and the like.

Preferred examples of substituted -A<sup>4</sup>-R<sub>d</sub> include methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, pentyl group, isoamyl group, cyclopropyl group, cyclopropylmethyl group, 2-(cyclopropyl)ethyl group, cyclopentyl group, cyclopentylmethyl group, 2-(cyclopentyl)ethyl group, cyclohexyl group, cyclohexylmethyl group, 2-(cyclohexyl)ethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, 2-(4-chlorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, (pyridin-2-yl)methyl group, (pyridin-3-yl)methyl group, (pyridin-4-yl)methyl group, and the like.

Particularly preferred examples of substituted -A<sup>4</sup>-R<sub>d</sub> include methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, pentyl group, isoamyl group, cyclopropyl group, cyclopropylmethyl group, cyclopentyl group, cyclopentylmethyl group, cyclohexyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, 2-(4-chlorophenyl)ethyl

group, 2-(4-fluorophenyl)ethyl group, and the like.

Preferred examples of Re include methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, phenylmethyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, furan-2-yl group, furan-3-yl group, thiophen-2-yl group, thiophen-3-yl group, methoxy group, ethoxy group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenoxy group, 4-methylphenoxy group, 4-chlorophenoxy group, 4-fluorophenoxy group, methylthio group, amino group, N-methylamino group, N,N-dimethylamino group, N-ethylamino group, N,N-diethylamino group, N-propylamino group, N-isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group, N-(4-fluorophenyl)amino group, N-(pyridin-2-yl)amino group, N-(pyridin-3-yl)amino group, N-(pyridin-4-yl)amino group, N-(furan-2-yl)amino group, N-(furan-3-yl)amino group, N-(thiophen-2-yl)amino group, N-(thiophen-3-yl)amino group, pyrrolidino group, piperidino group, morpholino group, methyloxycarbonylamino group, ethyloxycarbonylamino group, and the like.

Particularly preferred examples of Re include isopropyl group, butyl group, isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, propyloxy group, isopropyloxy

group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenyloxy group, 4-methylphenyloxy group, 4-chlorophenyloxy group, 4-fluorophenyloxy group, N-propylamino group, N-isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group, N-(4-fluorophenyl)amino group, pyrrolidino group, piperidino group, morpholino group, and the like.

Preferred examples of  $A^5-Re$  include acetyl group, thioacetyl group, methanesulfonyl group, propionyl group, ethylthiocarbonyl group, butyryl group, propylthiocarbonyl group, isobutyryl group, isopropylthiocarbonyl group, isopropylsulfonyl group, valeryl group, butylthiocarbonyl group, isovaleryl group, isobutylthiocarbonyl group, pivaloyl group, t-butylthiocarbonyl group, cyclopropylcarbonyl group, cyclopropylthiocarbonyl group, cyclopentylcarbonyl group, cyclopentylthiocarbonyl group, cyclohexylcarbonyl group, cyclohexylthiocarbonyl group, cyclopentylmethylcarbonyl group, cyclopentylmethylthiocarbonyl group, cyclohexylmethylcarbonyl group, cyclohexylmethylthiocarbonyl group, benzoyl group, thiobenzoyl group, phenylsulfonyl group, 4-methylphenylcarbonyl group, 4-methylphenylthiocarbonyl group, 4-methylphenylsulfonyl group, 4-chlorophenylcarbonyl group, 4-chlorophenylthiocarbonyl group, 4-fluorophenylcarbonyl group, 4-fluorophenylthiocarbonyl group, phenylmethylcarbonyl group, 4-methylphenylmethylcarbonyl group, 4-chlorophenylmethylcarbonyl group, 4-fluorophenylmethylcarbonyl group, (pyridin-2-yl)carbonyl group, (pyridin-2-yl)thiocarbonyl group, (pyridin-3-yl)carbonyl group, (pyridin-4-yl)carbonyl group, (furan-2-yl)carbonyl group, (thiophen-2-yl)carbonyl

group, methyloxycarbonyl group, methylsulfanylcabonyl group,  
 methoxythiocarbonyl group, methyloxycarbonylaminocarbonyl group, carbamoyl  
 group, N-methylcarbamoyl group, N-methylthiocarbamoyl group,  
 N,N-dimethylcarbamoyl group, N,N-dimethylthiocarbamoyl group,  
 N,N-dimethylsulfamoyl group, ethyloxycarbonyl group,  
 ethyloxycarbonylaminocarbonyl group, N-ethylcarbamoyl group,  
 N-ethylthiocarbamoyl group, N,N-diethylcarbamoyl group,  
 N,N-diethylthiocarbamoyl group, N,N-diethylsulfamoyl group, propyloxycarbonyl  
 group, N-propylcarbamoyl group, N-propylthiocarbamoyl group,  
 isopropyloxycarbonyl group, N-isopropylcarbamoyl group, N-isopropylthiocarbamoyl  
 group, butyloxycarbonyl group, N-butylcarbamoyl group, N-butylthiocarbamoyl  
 group, isobutyloxycarbonyl group, N-isobutylcarbamoyl group,  
 N-isobutylthiocarbamoyl group, t-butyloxycarbonyl group, N-t-butylcarbamoyl  
 group, N-t-butylthiocarbamoyl group, cyclopropyloxycarbonyl group,  
 N-cyclopropylcarbamoyl group, N-cyclopropylthiocarbamoyl group,  
 cyclopentyloxycarbonyl group, N-cyclopentylcarbamoyl group,  
 N-cyclopentylthiocarbamoyl group, cyclohexyloxycarbonyl group,  
 N-cyclohexylcarbamoyl group, N-cyclohexylthiocarbamoyl group,  
 cyclopentylmethyloxycarbonyl group, cyclohexylmethyloxycarbonyl group,  
 phenyloxycarbonyl group, N-phenylcarbamoyl group, N-phenylthiocarbamoyl group,  
 4-methylphenyloxycarbonyl group, N-(4-methylphenyl)carbamoyl group,  
 N-(4-methylphenyl)thiocarbamoyl group, 4-chlorophenyloxycarbonyl group,  
 N-(4-chlorophenyl)carbamoyl group, N-(4-chlorophenyl)thiocarbamoyl group,  
 4-fluorophenyloxycarbonyl group, N-(4-fluorophenyl)carbamoyl group,  
 N-(4-fluorophenyl)thiocarbamoyl group, phenylmethyloxycarbonyl group,  
 4-methylphenylmethyloxycarbonyl group, 4-chlorophenylmethyloxycarbonyl group,  
 4-fluorophenylmethyloxycarbonyl group, N-(pyridin-2-yl)carbamoyl group,

N-(pyridin-2-yl)thiocarbamoyl group, N-(pyridin-3-yl)carbamoyl group, N-(pyridin-3-yl)thiocarbamoyl group, N-(pyridin-4-yl)carbamoyl group, N-(pyridin-4-yl)thiocarbamoyl group, N-(furan-2-yl)carbamoyl group, N-(thiophen-2-yl)carbamoyl group, (pyrrolidino-1-yl)carbonyl group, (piperidino-1-yl)carbonyl group, (morpholino-4-yl)carbonyl group, and the like.

Particularly preferred examples of -A<sup>5</sup>-Re include isobutyryl group, isopropylthiocarbonyl group, isopropylsulfonyl group, valeryl group, butylthiocarbonyl group, isovaleryl group, isobutylthiocarbonyl group, pivaloyl group, t-butylthiocarbonyl group, cyclopropylcarbonyl group, cyclopropylthiocarbonyl group, cyclopentylcarbonyl group, cyclopentylthiocarbonyl group, cyclohexylcarbonyl group, cyclohexylthiocarbonyl group, cyclopentylmethylcarbonyl group, cyclopentylmethylthiocarbonyl group, cyclohexylmethylcarbonyl group, cyclohexylmethylthiocarbonyl group, benzoyl group, thiobenzoyl group, phenylsulfonyl group, 4-methylphenylcarbonyl group, 4-methylphenylthiocarbonyl group, 4-methylphenylsulfonyl group, 4-chlorophenylcarbonyl group, 4-chlorophenylthiocarbonyl group, 4-fluorophenylcarbonyl group, 4-fluorophenylthiocarbonyl group, isopropoxy carbonyl group, N-isopropylcarbamoyl group, N-isopropylthiocarbamoyl group, butyloxy carbonyl group, N-butylcarbamoyl group, N-butylthiocarbamoyl group, isobutyloxy carbonyl group, N-isobutylcarbamoyl group, N-isobutylthiocarbamoyl group, t-butyloxy carbonyl group, N-t-butylcarbamoyl group, N-t-butylthiocarbamoyl group, cyclopropyloxy carbonyl group, N-cyclopropylcarbamoyl group, N-cyclopropylthiocarbamoyl group, cyclopentyloxy carbonyl group, N-cyclopentylcarbamoyl group, N-cyclopentylthiocarbamoyl group, cyclohexyloxy carbonyl group, N-cyclohexylcarbamoyl group, N-cyclohexylthiocarbamoyl group, cyclopentylmethyloxy carbonyl group, cyclohexylmethyloxy carbonyl group,

phenyloxycarbonyl group, N-phenylcarbamoyl group, N-phenylthiocarbamoyl group, 4-methylphenyloxycarbonyl group, N-(4-methylphenyl)carbamoyl group, N-(4-methylphenyl)thiocarbamoyl group, 4-chlorophenyloxycarbonyl group, N-(4-chlorophenyl)carbamoyl group, N-(4-chlorophenyl)thiocarbamoyl group, 4-fluorophenyloxycarbonyl group, N-(4-fluorophenyl)carbamoyl group, N-(4-fluorophenyl)thiocarbamoyl group, (pyrrolidino-1-yl)carbonyl group, (piperidino-1-yl)carbonyl group, (morpholino-4-yl)carbonyl group, and the like.

Specific examples of R<sub>c</sub> include 2-(N-isobutyryl-N-methylamino)ethyl group, 2-(N-ethyl-N-isobutyrylamino)ethyl group, 2-(N-isobutyryl-N-propylamino)ethyl group, 2-(N-isobutyryl-N-isopropylamino)ethyl group, 2-(N-butyl-N-isobutyrylamino)ethyl group, 2-(N-isobutyl-N-isobutyrylamino)ethyl group, 2-(N-cyclopropyl-N-isobutyrylamino)ethyl group, 2-(N-cyclopentyl-N-isobutyrylamino)ethyl group, 2-(N-cyclopentylmethyl-N-isobutyrylamino)ethyl group, 2-(N-cyclohexyl-N-isobutyrylamino)ethyl group, 2-(N-cyclohexylmethyl-N-isobutyrylamino)ethyl group, 2-(N-isobutyryl-N-phenylamino)ethyl group, 2-[N-isobutyryl-N-(4-methylphenyl)amino]ethyl group, 2-[N-(4-chlorophenyl)-N-isobutyrylamino]ethyl group, 2-[N-(4-fluorophenyl)-N-isobutyrylamino]ethyl group, 2-(N-benzyl-N-isobutyrylamino)ethyl group, 2-[N-(4-chlorophenylmethyl)-N-isobutyrylamino]ethyl group, 2-[N-(4-fluorophenylmethyl)-N-isobutyrylamino]ethyl group, 2-[N-[2-(4-chlorophenyl)ethyl]-N-isobutyrylamino]ethyl group, 2-[N-[2-(4-fluorophenyl)ethyl]-N-isobutyrylamino]ethyl group, 2-(N-isobutylthiocarbonyl-N-methylamino)ethyl group, 2-(N-isobutylthiocarbonyl-N-isopropylamino)ethyl group,

2-(N-butyl-N-isobutylthiocarbonylamino)ethyl group,  
 2-(N-isobutyl-N-isobutylthiocarbonylamino)ethyl group,  
 2-(N-cyclopentyl-N-isobutylthiocarbonylamino)ethyl group,  
 2-(N-cyclopentylmethyl-N-isobutylthiocarbonylamino)ethyl group,  
 2-(N-isobutylthiocarbonyl-N-phenylamino)ethyl group,  
 2-(N-benzyl-N-isobutylthiocarbonylamino)ethyl group,  
 2-[N-(4-fluorophenylmethyl)-N-isobutylthiocarbonylamino]ethyl group,  
 2-(N-methyl-N-pivaloylamino)ethyl group, 2-(N-isopropyl-N-pivaloylamino)ethyl  
 group, 2-(N-butyl-N-pivaloylamino)ethyl group, 2-(N-isobutyl-N-pivaloylamino)ethyl  
 group, 2-(N-cyclohexyl-N-pivaloylamino)ethyl group,  
 2-(N-cyclohexylmethyl-N-pivaloylamino)ethyl group,  
 2-(N-phenyl-N-pivaloylamino)ethyl group, 2-(N-benzyl-N-pivaloylamino)ethyl group,  
 2-(N-cyclopentylcarbonyl-N-methylamino)ethyl group,  
 2-(N-butyl-N-cyclopentylcarbonylamino)ethyl group,  
 2-(N-cyclopentylcarbonyl-N-isobutylamino)ethyl group,  
 2-(N-cyclopentylcarbonyl-N-cyclopentylmethylamino)ethyl group,  
 2-(N-cyclopentylcarbonyl-N-phenylamino)ethyl group,  
 2-[N-cyclopentylcarbonyl-N-(4-fluorophenyl)amino]ethyl group,  
 2-(N-benzyl-N-cyclopentylcarbonylamino)ethyl group,  
 2-[N-cyclopentylcarbonyl-N-(4-fluorophenylmethyl)amino]ethyl group,  
 2-(N-methyl-N-phenylsulfonylamino)ethyl group,  
 2-(N-ethyl-N-phenylsulfonylamino)ethyl group,  
 2-(N-phenylsulfonyl-N-propylamino)ethyl group,  
 2-(N-isopropyl-N-phenylsulfonylamino)ethyl group,  
 2-(N-butyl-N-phenylsulfonylamino)ethyl group,  
 2-(N-isobutyl-N-phenylsulfonylamino)ethyl group,  
 2-(N-cyclopropyl-N-phenylsulfonylamino)ethyl group,



2-(N-cyclopentyl-N-phenylsulfonylamino)ethyl group,  
 2-(N-cyclopentylmethyl-N-phenylsulfonylamino)ethyl group,  
 2-(N-cyclohexyl-N-phenylsulfonylamino)ethyl group,  
 2-(N-cyclohexylmethyl-N-phenylsulfonylamino)ethyl group,  
 2-(N-phenyl-N-phenylsulfonylamino)ethyl group,  
 2-[N-(4-fluorophenyl)-N-phenylsulfonylamino]ethyl group,  
 2-(N-benzyl-N-phenylsulfonylamino)ethyl group,  
 2-[N-(N-butylcarbamoyl)-N-methylamino]ethyl group,  
 2-[N-butyl-N-(N-butylcarbamoyl)amino]ethyl group,  
 2-[N-(N-butylcarbamoyl)-N-isobutylamino]ethyl group,  
 2-[N-(N-butylcarbamoyl)-N-cyclopentylamino]ethyl group,  
 2-[N-(N-butylcarbamoyl)-N-cyclohexylmethylamino]ethyl group,  
 2-[N-(N-butylcarbamoyl)-N-phenylamino]ethyl group,  
 2-{N-(N-butylcarbamoyl)-N-(4-fluorophenyl)amino}ethyl group,  
 2-[N-benzyl-N-(N-butylcarbamoyl)amino]ethyl group,  
 2-{N-(N-butylcarbamoyl)-N-(4-fluorophenylmethyl)amino}ethyl group,  
 2-{N-(N-butylcarbamoyl)-N-[2-(4-fluorophenyl)ethyl]amino}ethyl group,  
 2-[N-(N-isopropylthiocarbamoyl)-N-methylamino]ethyl group,  
 2-[N-butyl-N-(N-isopropylthiocarbamoyl)amino]ethyl group,  
 2-[N-isobutyl-N-(N-isopropylthiocarbamoyl)amino]ethyl group,  
 2-[N-cyclopentyl-N-(N-isopropylthiocarbamoyl)amino]ethyl group,  
 2-[N-cyclohexylmethyl-N-(N-isopropylthiocarbamoyl)amino]ethyl group,  
 2-[N-(N-isopropylthiocarbamoyl)-N-phenylamino]ethyl group,  
 2-{N-(4-fluorophenyl)-N-(N-isopropylthiocarbamoyl)amino}ethyl group,  
 2-[N-benzyl-N-(N-isopropylthiocarbamoyl)amino]ethyl group,  
 2-(N-isobutyloxycarbonyl-N-methylamino)ethyl group,  
 2-(N-butyl-N-isobutyloxycarbonylamino)ethyl group,

2-(N-isobutyl-N-isobutyloxycarbonylamino)ethyl group,  
 2-(N-cyclopentyl-N-isobutyloxycarbonylamino)ethyl group,  
 2-(N-cyclohexylmethyl-N-isobutyloxycarbonylamino)ethyl group,  
 2-(N-isobutyloxycarbonyl-N-phenylamino)ethyl group,  
 2-[N-(4-fluorophenyl)-N-isobutyloxycarbonylamino]ethyl group,  
 2-(N-benzyl-N-isobutyloxycarbonylamino)ethyl group,  
 2-[N-(N-cyclopentylcarbamoyl)-N-methylamino]ethyl group,  
 2-[N-butyl-N-(N-cyclopentylcarbamoyl)amino]ethyl group,  
 2-[N-(N-cyclopentylcarbamoyl)-N-isobutylamino]ethyl group,  
 2-[N-cyclopentyl-N-(N-cyclopentylcarbamoyl)amino]ethyl group,  
 2-[N-cyclohexylmethyl-N-(N-cyclopentylcarbamoyl)amino]ethyl group,  
 2-[N-(N-cyclopentylcarbamoyl)-N-phenylamino]ethyl group,  
 2-[N-benzyl-N-(N-cyclopentylcarbamoyl)amino]ethyl group,  
 2-[N-(N-cyclohexylthiocarbamoyl)-N-methylamino]ethyl group,  
 2-[N-butyl-N-(N-cyclohexylthiocarbamoyl)amino]ethyl group,  
 2-[N-(N-cyclohexylthiocarbamoyl)-N-isobutylamino]ethyl group,  
 2-[N-(N-cyclohexylthiocarbamoyl)-N-cyclopentylamino]ethyl group,  
 2-[N-cyclohexylmethyl-N-(N-cyclohexylthiocarbamoyl)amino]ethyl group,  
 2-[N-(N-cyclohexylthiocarbamoyl)-N-phenylamino]ethyl group,  
 2-[N-benzyl-N-(N-cyclohexylthiocarbamoyl)amino]ethyl group,  
 2-(N-methyl-N-phenyloxycarbonylamino)ethyl group,  
 2-(N-butyl-N-phenyloxycarbonylamino)ethyl group,  
 2-(N-isobutyl-N-phenyloxycarbonylamino)ethyl group,  
 2-(N-cyclopentyl-N-phenyloxycarbonylamino)ethyl group,  
 2-(N-cyclohexylmethyl-N-phenyloxycarbonylamino)ethyl group,  
 2-(N-phenyl-N-phenyloxycarbonylamino)ethyl group,  
 2-(N-benzyl-N-phenyloxycarbonylamino)ethyl group,

2-[N-methyl-N-(N-phenylcarbamoyl)amino]ethyl group,  
 2-[N-butyl-N-(N-phenylcarbamoyl)amino]ethyl group,  
 2-[N-isobutyl-N-(N-phenylcarbamoyl)amino]ethyl group,  
 2-[N-cyclopentyl-N-(N-phenylcarbamoyl)amino]ethyl group,  
 2-[N-cyclohexylmethyl-N-(N-phenylcarbamoyl)amino]ethyl group,  
 2-[N-phenyl-N-(N-phenylcarbamoyl)amino]ethyl group,  
 2-[N-benzyl-N-(N-phenylcarbamoyl)amino]ethyl group, and the like.

When Rs in the formula (I) represents  $\cdot N(R_y)(R_z)$ ,  $R_z$  is defined to have the same meaning as  $R_x$ , or  $R_z$  represents methyl group, ethyl group, or a  $\cdot A^5\text{-Re}$  group.  $\cdot A^5\text{-Re}$  has the same meaning as defined above.

Particularly preferred examples of  $R_z$  include butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl

group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group,  
 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group,  
 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group,  
 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group,  
 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group,  
 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group,  
 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group,  
 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group,  
 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group,  
 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group,  
 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group,  
 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group,  
 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,  
 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,  
 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group,  
 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group,  
 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group,  
 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group,  
 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group,  
 2-(N-phenyl-N-methylamino)ethyl group, 2-(N-ethyl-N-phenylamino)ethyl group,  
 isobutyryl group, isopropylthiocarbonyl group, isopropylsulfonyl group, valeryl  
 group, butylthiocarbonyl group, isovaleryl group, isobutylthiocarbonyl group,  
 pivaloyl group, t-butylthiocarbonyl group, cyclopropylcarbonyl group,  
 cyclopropylthiocarbonyl group, cyclopentylcarbonyl group, cyclopentylthiocarbonyl  
 group, cyclohexylcarbonyl group, cyclohexylthiocarbonyl group,  
 cyclopentylmethylcarbonyl group, cyclopentylmethylthiocarbonyl group,  
 cyclohexylmethylcarbonyl group, cyclohexylmethylthiocarbonyl group, benzoyl

group, thiobenzoyl group, phenylsulfonyl group, 4-methylphenylcarbonyl group, 4-methylphenylthiocarbonyl group, 4-methylphenylsulfonyl group, 4-chlorophenylcarbonyl group, 4-chlorophenylthiocarbonyl group, 4-fluorophenylcarbonyl group, 4-fluorophenylthiocarbonyl group, isopropoxy carbonyl group, N-isopropyl carbamoyl group, N-isopropylthiocarbamoyl group, butyloxy carbonyl group, N-butyl carbamoyl group, N-butylthiocarbamoyl group, isobutyloxy carbonyl group, N-isobutyl carbamoyl group, N-isobutylthiocarbamoyl group, t-butyloxy carbonyl group, N-t-butyl carbamoyl group, N-t-butylthiocarbamoyl group, cyclopropyloxy carbonyl group, N-cyclopropyl carbamoyl group, N-cyclopropylthiocarbamoyl group, cyclopentyloxy carbonyl group, N-cyclopentyl carbamoyl group, N-cyclopentylthiocarbamoyl group, cyclohexyloxy carbonyl group, N-cyclohexyl carbamoyl group, N-cyclohexylthiocarbamoyl group, cyclopentylmethyloxy carbonyl group, cyclohexylmethyloxy carbonyl group, phenyloxy carbonyl group, N-phenyl carbamoyl group, N-phenylthiocarbamoyl group, 4-methylphenyloxy carbonyl group, N-(4-methylphenyl) carbamoyl group, N-(4-methylphenyl)thiocarbamoyl group, 4-chlorophenyloxy carbonyl group, N-(4-chlorophenyl) carbamoyl group, N-(4-chlorophenyl)thiocarbamoyl group, 4-fluorophenyloxy carbonyl group, N-(4-fluorophenyl) carbamoyl group, N-(4-fluorophenyl)thiocarbamoyl group, (pyrrolidino-1-yl) carbonyl group, (piperidino-1-yl) carbonyl group, (morpholino-4-yl) carbonyl group, and the like.

Among the R<sub>z</sub>, methyl group or ethyl group is particularly preferred when R<sub>y</sub> is other than hydrogen atom.

R<sub>y</sub> represents hydrogen atom, an alkyl group having 1 to 8 carbon atoms, or a -A<sup>6</sup>-Q<sub>p</sub> group, or binds to R<sub>z</sub> to form a saturated or unsaturated nitrogen-containing cyclic substituent having 3 to 7 atoms together with nitrogen atom to which they bind. The alkyl group having 1 to 8 carbon atoms is a linear or

branched saturated alkyl group, a linear or branched partially unsaturated alkyl group, or an alkyl group which may contain a cyclic alkyl group having 3 to 7 carbon atoms. Examples include methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, pentyl group, isopentyl group, 2-methylbutyl group, 2,2-dimethylpropyl group, hexyl group, 4-methylpentyl group, 2,3-dimethylbutyl group, 2-ethylbutyl group, heptyl group, octyl group, cyclopropyl group, cyclobutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopropylmethyl group, cyclobutylmethyl group, cyclopentylmethyl group, cyclohexylmethyl group, cycloheptylmethyl group, 2-cyclopentylethyl group, 2-cyclohexylethyl group, 2-methylcyclopentyl group, 3-methylcyclopentyl group, 3,4-dimethylcyclopentyl group, 4-methylcyclohexyl group, 4,4-dimethylcyclohexyl group, 4-ethylcyclohexyl group, 4-methylcyclohexylmethyl group, and the like.

-A<sup>6</sup>-Qp has the same meaning as defined above.

Particularly preferred examples of Ry include hydrogen atom, methyl group, ethyl group, isobutyl group, and the like.

Ry also binds to Rz to represents a saturated or unsaturated nitrogen-containing cyclic substituent having 3 to 7 atoms formed together with the nitrogen atom to which they bind. Specific examples thereof include cyclic substituents containing nitrogen atom such as 1-pyrrolidino group, 1-piperidino group, 1-homopiperidino group, 1-piperazino group, 4-morpholino group, pyrrol-1-yl group, imidazol-1-yl group, and pyrazol-1-yl group, and all of these are preferred.

The nitrogen-containing cyclic substituent may be substituted with one or two lower alkyl groups having 1 to 4 carbon atoms wherein the two alkyl groups may be the same or different. Examples of the lower alkyl having 1 to 4 carbon atoms include methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, or t-butyl group.

Among the substituent -N(Ry)(Rz), particularly preferred examples include

N,N-dimethylamino group, N-ethyl-N-methylamino group, N,N-diethylamino group,  
 N-methyl-N-propylamino group, N-ethyl-N-propylamino group,  
 N-isopropyl-N-methylamino group, N-ethyl-N-isopropylamino group, N-butylamino  
 group, N-butyl-N-methylamino group, N-butyl-N-ethylamino group,  
 N-isobutylamino group, N-isobutyl-N-methylamino group, N-ethyl-N-isobutylamino  
 group, N-(2-ethylbutyl)amino group, N-(2-ethylbutyl)-N-methylamino group,  
 N-cyclopentylamino group, N-cyclopentyl-N-methylamino group, N-cyclohexylamino  
 group, N-cyclohexyl-N-methylamino group, N-cycloheptylamino group,  
 N-(cyclopentylmethyl)amino group, N-(cyclopentylmethyl)-N-methylamino group,  
 N-(cyclohexylmethyl)amino group, N-(cyclohexylmethyl)-N-methylamino group,  
 N-(2-methylphenyl)amino group, N-(4-methylphenyl)amino group,  
 N-(2-fluorophenyl)amino group, N-(3-fluorophenyl)amino group,  
 N-(4-fluorophenyl)amino group, N-(2-chlorophenyl)amino group,  
 N-(3-chlorophenyl)amino group, N-(4-chlorophenyl)amino group,  
 N-(indan-2-yl)amino group, N-(1-phenylethyl)amino group,  
 N-[1-(2-fluorophenyl)ethyl]amino group, N-[1-(3-fluorophenyl)ethyl]amino group,  
 N-[1-(4-fluorophenyl)ethyl]amino group, N-[1-(2-chlorophenyl)ethyl]amino group,  
 N-[1-(3-chlorophenyl)ethyl]amino group, N-[1-(4-chlorophenyl)ethyl]amino group,  
 N-(2-methylphenylmethyl)amino group, N-methyl-N-(2-methylphenylmethyl)amino  
 group, N-(3-methylphenylmethyl)amino group,  
 N-methyl-N-(3-methylphenylmethyl)amino group, N-(4-methylphenylmethyl)amino  
 group, N-methyl-N-(4-methylphenylmethyl)amino group,  
 N-(2-fluorophenylmethyl)amino group, N-(2-fluorophenylmethyl)-N-methylamino  
 group, N-(3-fluorophenylmethyl)amino group,  
 N-(3-fluorophenylmethyl)-N-methylamino group, N-(4-fluorophenylmethyl)amino  
 group, N-(4-fluorophenylmethyl)-N-methylamino group,  
 N-(2-chlorophenylmethyl)amino group, N-(2-chlorophenylmethyl)-N-methylamino

group, N-(3-chlorophenylmethyl)amino group,  
 N-(3-chlorophenylmethyl)-N-methylamino group, N-(4-chlorophenylmethyl)amino  
 group, N-(4-chlorophenylmethyl)-N-methylamino group,  
 N-(2,3-difluorophenylmethyl)amino group,  
 N-(2,3-difluorophenylmethyl)-N-methylamino group,  
 N-(2,4-difluorophenylmethyl)amino group,  
 N-(2,4-difluorophenylmethyl)-N-methylamino group,  
 N-(2,5-difluorophenylmethyl)amino group,  
 N-(2,5-difluorophenylmethyl)-N-methylamino group,  
 N-(3,4-difluorophenylmethyl)amino group,  
 N-(3,4-difluorophenylmethyl)-N-methylamino group,  
 N-(3,5-difluorophenylmethyl)amino group,  
 N-(3,5-difluorophenylmethyl)-N-methylamino group,  
 N-(2,3-dichlorophenylmethyl)amino group,  
 N-(2,3-dichlorophenylmethyl)-N-methylamino group,  
 N-(2,4-dichlorophenylmethyl)amino group,  
 N-(2,4-dichlorophenylmethyl)-N-methylamino group,  
 N-(2,5-dichlorophenylmethyl)amino group,  
 N-(2,5-dichlorophenylmethyl)-N-methylamino group,  
 N-(2,6-dichlorophenylmethyl)amino group,  
 N-(2,6-dichlorophenylmethyl)-N-methylamino group,  
 N-(3,4-dichlorophenylmethyl)amino group,  
 N-(3,4-dichlorophenylmethyl)-N-methylamino group,  
 N-(3,5-dichlorophenylmethyl)amino group,  
 N-(3,5-dichlorophenylmethyl)-N-methylamino group,  
 N-[2-(trifluoromethyl)phenylmethyl]amino group,  
 N-methyl-N-[2-(trifluoromethyl)phenylmethyl]amino group,



N-[3-(trifluoromethyl)phenylmethyl]amino group,  
 N-methyl-N-[3-(trifluoromethyl)phenylmethyl]amino group,  
 N-[4-(trifluoromethyl)phenylmethyl]amino group,  
 N-methyl-N-[4-(trifluoromethyl)phenylmethyl]amino group, 1-pyrrolidino group,  
 1-(4-methylpiperidino) group, 1-homopiperidino group, and 4-morpholino group.

A most preferred example of Rs in the aforementioned general formula (I) include Rs which meets the conditions of: Rs is  $\cdot D \cdot R_x$  wherein D is a single bond and  $R_x$  represents Rb, and  $A^1$  and  $A^2$  in Rb are single bonds. Specific examples include phenyl group, 2-methylphenyl group, 3-methylphenyl group, 4-methylphenyl group, 2,3-dimethylphenyl group, 3,5-dimethylphenyl group, 2-methoxyphenyl group, 3-methoxyphenyl group, 4-methoxyphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, 2,3-difluorophenyl group, 2,4-difluorophenyl group, 2,5-difluorophenyl group, 3,4-difluorophenyl group, 2,3-dichlorophenyl group, 2,4-dichlorophenyl group, 2,5-dichlorophenyl group, 2,6-dichlorophenyl group, 3,4-dichlorophenyl group, 3,5-dichlorophenyl group, 2-trifluoromethylphenyl group, 3-trifluoromethylphenyl group, 4-trifluoromethylphenyl group, 4-(N,N-dimethylamino)phenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, furan-2-yl group, furan-3-yl group, thiophen-2-yl group, thiophen-3-yl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, naphthalen-1-yl group, naphthalen-2-yl group, 1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1H-indazol-5-yl group, or

1-methyl-1H-indazol-5-yl group.

AR in the formula (I) is defined to be a residue of a partially unsaturated or completely unsaturated condensed bicyclic carbon ring or heterocyclic ring (ar).

Further, AR may be substituted with one of Xa or two or more of the same or different Xa. The heterocyclic ring (ar) means a ring containing 1 to 4 the same or different ring-constituting heteroatoms selected from the group consisting of nitrogen atom, oxygen atom, and sulfur atom.

The "condensed bicyclic carbon ring or heterocyclic ring" means a partially unsaturated or completely unsaturated ring having 8 to 11 atoms. Preferred examples include a partially unsaturated or completely unsaturated ring consisting of 8 atoms formed by fusion of 5-membered heterocyclic rings containing 1 or 2 ring-constituting heteroatoms selected from the group consisting of nitrogen, oxygen, and sulfur atoms, a partially unsaturated or completely unsaturated ring consisting of 9 atoms formed by fusion of a 5-membered heterocyclic ring containing 1 or 2 ring-constituting heteroatoms selected from the group consisting of nitrogen, oxygen, and sulfur atoms and a 6-membered carbon ring or a 6-membered heterocyclic ring containing 1 or 2 ring-constituting heteroatoms selected from the group consisting of nitrogen, oxygen, and sulfur atoms, and a partially unsaturated or completely unsaturated substituent consisting of 10 atoms formed by fusion of a 6-membered carbon ring or a 6-membered heterocyclic ring containing 1 or 2 ring-constituting heteroatoms selected from the group consisting of nitrogen, oxygen, and sulfur atoms and a 6-membered carbon ring or 6-membered heterocyclic rings containing 1 or 2 ring-constituting heteroatoms selected from the group consisting of nitrogen, oxygen, and sulfur atom. As the carbon ring constituting AR not containing a heteroatom, among the rings constituting AR, naphthalene ring is particularly preferred. Further, as the heterocyclic ring (ar) containing a heteroatom, among the rings constituting AR, those containing 1 or 2 ring-constituting heteroatoms are

preferred.

As for AR in the formula (I), specific examples of preferred ring constituting AR include naphthalene, benzofuran, benzo[b]thiophene, indole, benzothiazole, dihydro-3H-benzothiazole, quinoline, dihydro-1H-quinoline, benzo[d]isothiazole, 1H-indazole, benzo[c]isothiazole, 2H-indazole, imidazo[1,2-a]pyridine, 1H-pyrrolo[2,3-b]pyridine, isoquinoline, dihydro-2H-isoquinoline, cinnoline, quinazoline, quinoxaline, 1H-benzimidazole, benzoxazole, 1H-pyrrolo[3,2-b]pyridine, benzo[1,2,5]thiadiazole, 1H-benzotriazole, 1,3-dihydropyrrolo[2,3-b]pyridine, 1,3-dihydrobenzimidazole, dihydro-3H-benzoxazole, phthalazine, [1,8]naphthalidine, [1,5]naphthalidine, 1H-pyrrolo[3,2-c]pyridine, 1H-pyrrolo[2,3-c]pyridine, 1H-pyrazolo[4,3-b]pyridine, 1H-pyrazolo[4,3-c]pyridine, 1H-pyrazolo[3,4-c]pyridine, 1H-pyrazolo[3,4-b]pyridine, [1,2,4]triazolo[4,3-a]pyridine, thieno[3,2-c]pyridine, thieno[3,2-b]pyridine, 1H-thieno[3,2-c]pyrazole, benzo[d]isoxazole, benzo[c]isoxazole, indolizine, 1,3-dihydroindol, 1H-pyrazolo[3,4-d]thiazole, 2H-isoindol, [1,2,4]triazolo[1,5-a]pyrimidine, 1H-pyrazolo[3,4-b]pyrazine, 1H-imidazo[4,5-b]pyrazine, 7H-purine, 4H-chromene, and the like. Among them, naphthalene, benzofuran, benzo[b]thiophene, indole, benzothiazole, dihydro-3H-benzothiazole, quinoline, dihydro-1H-quinoline, benzo[d]isothiazole, 1H-indazole, benzo[c]isothiazole, 2H-indazole, imidazo[1,2-a]pyridine, 1H-pyrrolo[2,3-b]pyridine, isoquinoline and dihydro-2H-isoquinoline constitute a particularly preferred group, and cinnoline, quinazoline, quinoxaline, 1H-benzimidazole, benzoxazole, 1H-pyrrolo[3,2-b]pyridine, benzo[1,2,5]thiadiazole, 1H-benzotriazole, 1,3-dihydropyrrolo[2,3-b]pyridine, 1,3-dihydrobenzimidazole and dihydro-3H-benzoxazole also constitute a particularly preferred group. Further, naphthalene, benzofuran, benzo[b]thiophene, indole, benzothiazole, quinoline, 1H-indazole and isoquinoline are particularly preferred.

AR binds to any of the ring-constituting carbon atoms C<sup>2</sup>, C<sup>3</sup>, C<sup>4</sup>, C<sup>5</sup>, and C<sup>6</sup>

in the aromatic ring (E) in the aforementioned formula (I) at an arbitrary carbon atom in AR. Preferred examples of the ring constituting AR include, as indicated with substitution position in the aromatic ring (E), naphthalen-2-yl group, naphthalen-1-yl group, benzofuran-5-yl group, benzofuran-4-yl group, benzofuran-2-yl group, benzo[b]thiophen-5-yl group, benzo[b]thiophen-4-yl group, benzo[b]thiophen-2-yl group, indol-5-yl group, indol-4-yl group, indol-6-yl group, benzothiazol-6-yl group, benzothiazol-7-yl group, benzothiazol-5-yl group, benzothiazol-4-yl group, dihydro-3H-benzothiazol-6-yl group, dihydro-3H-benzothiazol-7-yl group, dihydro-3H-benzothiazol-5-yl group, dihydro-3H-benzothiazol-4-yl group, quinolin-6-yl group, quinolin-3-yl group, quinolin-5-yl group, quinolin-7-yl group, dihydro-1H-quinolin-6-yl group, dihydro-1H-quinolin-5-yl group, benzo[d]isothiazol-5-yl group, benzo[d]isothiazol-4-yl group, benzo[d]isothiazol-6-yl group, benzo[d]isothiazol-7-yl group, 1H-indazol-5-yl group, 1H-indazol-4-yl group, 1H-indazol-6-yl group, benzo[c]isothiazol-5-yl group, benzo[c]isothiazol-4-yl group, benzo[c]isothiazol-6-yl group, benzo[c]isothiazol-7-yl group, 2H-indazol-5-yl group, 2H-indazol-4-yl group, 2H-indazol-6-yl group, imidazo[1,2-a]pyridin-6-yl group, imidazo[1,2-a]pyridin-7-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1H-pyrrolo[2,3-b]pyridin-4-yl group, isoquinolin-6-yl group, isoquinolin-3-yl group, isoquinolin-5-yl group, isoquinolin-7-yl group, dihydro-2H-isoquinolin-6-yl group, dihydro-2H-isoquinolin-5-yl group, cinnolin-6-yl group, cinnolin-5-yl group, quinazolin-6-yl group, quinazolin-7-yl group, quinazolin-5-yl group, quinoxalin-2-yl group, quinoxalin-6-yl group, quinoxalin-5-yl group, 1H-benzimidazol-5-yl group, 1H-benzimidazol-4-yl group, benzoxazol-5-yl group, benzoxazol-6-yl group, benzoxazol-4-yl group, benzoxazol-7-yl group, 1H-pyrrolo[3,2-b]pyridin-5-yl group, 1H-pyrrolo[3,2-b]pyridin-6-yl group, benzo[1,2,5]thiadiazol-5-yl group, benzo[1,2,5]thiadiazol-4-yl group, 1H-benzotriazol-5-yl group, 1H-benzotriazol-4-yl

group, 1,3-dihydropyrrolo[2,3-b]pyridin-5-yl group,  
 1,3-dihydropyrrolo[2,3-b]pyridin-4-yl group, 1,3-dihydrobenzimidazol-5-yl group,  
 1,3-dihydrobenzimidazol-4-yl group, dihydro-3H-benzoxazol-6-yl group,  
 dihydro-3H-benzoxazol-7-yl group, dihydro-3H-benzoxazol-5-yl group,  
 dihydro-3H-benzoxazol-4-yl group, phthalazin-6-yl group, phthalazin-5-yl group,  
 [1,8]naphthalidin-3-yl group, [1,8]naphthalidin-4-yl group, [1,5]naphthalidin-3-yl  
 group, [1,5]naphthalidin-4-yl group, 1H-pyrrolo[3,2-c]pyridin-6-yl group,  
 1H-pyrrolo[3,2-c]pyridin-4-yl group, 1H-pyrrolo[2,3-c]pyridin-5-yl group,  
 1H-pyrrolo[2,3-c]pyridin-4-yl group, 1H-pyrazolo[4,3-b]pyridin-5-yl group,  
 1H-pyrazolo[4,3-b]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-6-yl group,  
 1H-pyrazolo[4,3-c]pyridin-4-yl group, 1H-pyrazolo[3,4-c]pyridin-5-yl group,  
 1H-pyrazolo[3,4-c]pyridin-4-yl group, 1H-pyrazolo[3,4-b]pyridin-5-yl group,  
 1H-pyrazolo[3,4-b]pyridin-4-yl group, [1,2,4]triazolo[4,3-a]pyridin-6-yl group,  
 [1,2,4]triazolo[4,3-a]pyridin-7-yl group, thieno[3,2-c]pyridin-2-yl group,  
 thieno[3,2-c]pyridin-3-yl group, thieno[3,2-c]pyridin-6-yl group,  
 thieno[3,2-b]pyridin-2-yl group, thieno[3,2-b]pyridin-3-yl group,  
 thieno[3,2-b]pyridin-5-yl group, thieno[3,2-b]pyridin-6-yl group,  
 1H-thieno[3,2-c]pyrazol-5-yl group, 1H-thieno[3,2-c]pyrazol-4-yl group,  
 benzo[d]isoxazol-5-yl group, benzo[d]isoxazol-4-yl group, benzo[d]isoxazol-6-yl group,  
 benzo[d]isoxazol-7-yl group, benzo[c]isoxazol-5-yl group, benzo[c]isoxazol-4-yl group,  
 benzo[c]isoxazol-6-yl group, benzo[c]isoxazol-7-yl group, indolizin-7-yl group,  
 indolizin-6-yl group, indolizine-8-yl group, 1,3-dihydroindol-5-yl group,  
 1,3-dihydroindol-4-yl group, 1,3-dihydroindol-6-yl group,  
 1H-pyrazolo[3,4-d]thiazol-5-yl group, 2H-isoindol-5-yl group, 2H-isoindol-4-yl group,  
 [1,2,4]triazolo[1,5-a]pyrimidin-6-yl group, 1H-pyrazolo[3,4-b]pyrazin-5-yl group,  
 1H-imidazo[4,5-b]pyrazin-5-yl group, 7H-purin-2-yl group, 4H-chromen-6-yl group,  
 4H-chromen-5-yl group, and the like. Among them, naphthalen-2-yl group,

naphthalen-1-yl group, benzofuran-5-yl group, benzofuran-4-yl group, benzo[b]thiophen-5-yl group, benzo[b]thiophen-4-yl group, indol-5-yl group, indol-4-yl group, benzothiazol-6-yl group, benzothiazol-7-yl group, quinolin-6-yl group, quinolin-3-yl group, dihydro-1H-quinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1H-indazol-4-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, dihydro-2H-isoquinolin-6-yl group, cinnolin-6-yl group, benzoxazol-5-yl group, and the like constitute a particularly preferred group, and naphthalen-2-yl group, benzofuran-5-yl group, benzo[b]thiophen-5-yl group, indol-5-yl group, benzothiazol-6-yl group, quinolin-6-yl group, quinolin-3-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, cinnolin-6-yl group, benzoxazol-5-yl group and the like are particularly preferred.

Further, AR may be substituted with one of Xa or the same or different two or more of Xa. Examples of substitution position of Xa include a carbon atom of AR not bonding to the aromatic ring (E), and/or when nitrogen atom is present, that nitrogen atom.

The substituent Xa represents a linear or branched saturated alkyl group having 1 to 4 carbon atoms, a saturated cyclic alkyl group having 3 to 7 carbon atoms, oxo group, thioxo group, fluorine atom, chlorine atom, trifluoromethyl group,  $-(CH_2)_iR^{14}$ ,  $-OR^{10}$ ,  $-N(R^{11})(R^{12})$ ,  $-SO_2R^{13}$ , or  $-COR^{27}$ . However, when nitrogen atom is present in AR, Xa which may substitute on the nitrogen atom represents a linear or branched saturated alkyl group having 1 to 4 carbon atoms, a saturated cyclic alkyl group having 3 to 7 carbon atoms, or  $-(CH_2)_iR^{14}$ .

Preferred examples of the substituent Xa are oxo group, thioxo group, fluorine atom, chlorine atom, and trifluoromethyl group.

Examples of the linear or branched saturated alkyl group having 1 to 4

carbon atoms as the substituent Xa include methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, and the like, and among them, methyl group, ethyl group, and propyl group are particularly preferred.

Further, examples of the saturated cyclic alkyl group having 3 to 7 carbon atoms include cyclopropyl group, cyclobutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, and the like.

$-(CH_2)_iR^{14}$  has the same meaning as defined above. Preferred examples are 2-hydroxyethyl group, carboxymethyl group, 2-carboxyethyl group, and N,N-dimethylcarbamoylmethyl group, and a particularly preferred example is 2-hydroxyethyl group.

$R^{10}$  in  $-OR^{10}$  represents hydrogen atom, a lower alkyl group having 1 to 4 carbon atoms, or a  $-(CH_2)_iR^{14}$  group, and among them, hydrogen atom is a particularly preferred example. Examples of the lower alkyl group having 1 to 4 carbon atoms include methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, and the like. Among them, methyl group is particularly preferred.  $-(CH_2)_iR^{14}$  has the same meaning as defined above. Therefore, preferred examples of  $-OR^{10}$  are hydroxyl group, methoxy group, 2-hydroxyethyloxy group, carboxymethyloxy group, 2-carboxyethyloxy group, N,N-dimethylcarbamoylmethyloxy group, and the like, and hydroxyl group, methoxy group, and 2-hydroxyethyloxy group are particularly preferred.

$R^{11}$  in  $-N(R^{11})(R^{12})$  represents hydrogen atom, or a lower alkyl group having 1 to 4 carbon atoms, and  $R^{12}$  represents hydrogen atom, a lower alkyl group having 1 to 4 carbon atoms, a hydroxyalkyl group having 2 to 4 carbon atoms,  $-COR^{15}$ , or  $-SO_2R^{16}$ , or binds to  $R^{11}$  to form a 3- to 6-membered ring together with the nitrogen atom to which they bind to form a saturated nitrogen-containing cycloalkyl group or morpholino group.  $R^{15}$  in  $-COR^{15}$  represents a lower alkyl group having 1 to 4 carbon atoms, a hydroxyalkyl group having 2 to 4 carbon atoms, amino group, a

mono- or dialkylamino group having 1 to 4 carbon atoms, or  $-A^6-Qa$ .  $R^{16}$  in  $-SO_2R^{16}$  represents a lower alkyl group having 1 to 4 carbon atoms, amino group, or a mono- or dialkylamino group having 1 to 4 carbon atoms. Specific examples of  $-N(R^{11})(R^{12})$  include amino group, N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, piperidino group, pyrrolidino group, morpholino group, 2-hydroxyethylamino group, formylamino group, acetylamino group, benzoyl group, furan-2-carboxyamino group, 2-hydroxyacetylamino group, 2-aminoacetylamino group, carbamoylamino group, N-methylcarbamoylamino group, N,N-dimethylcarbamoylamino group, methylsulfonylamino group, sulfamoylamino group, N-methylsulfamoylamino group, N,N-dimethylsulfamoylamino group, and the like. Among them, preferred examples are amino group, methylamino group, dimethylamino group, 2-hydroxyethylamino group, carbamoylamino group, acetylamino group, furan-2-carboxyamino group, 2-hydroxyacetylamino group, 2-aminoacetylamino group, methylsulfonylamino group, (N,N-dimethylsulfamoyl)amino group, and the like, and amino group, N-methylamino group, N,N-dimethylamino group, and 2-hydroxyethylamino group are particularly preferred.

$R^{13}$  in  $-SO_2R^{13}$  represents a lower alkyl group having 1 to 4 carbon atoms, amino group, or a mono- or dialkylamino group having 1 to 4 carbon atoms. Preferred examples of  $-SO_2R^{13}$  include methanesulfonyl group, sulfamoyl group, N-methylsulfamoyl group, N,N-dimethylsulfamoyl group, and the like.

$R^{27}$  in  $-COR^{27}$  represents hydrogen atom, hydroxyl group, an alkoxy group having 1 to 4 carbon atoms, a lower alkyl group having 1 to 4 carbon atoms, amino group, or a mono- or dialkylamino group having 1 to 4 carbon atoms. Specific examples of  $-COR^{27}$  include formyl group, carboxyl group, methoxycarbonyl group, ethoxycarbonyl group, acetyl group, propionyl group, carbamoyl group, N-methylcarbamoyl group, N,N-dimethylcarbamoyl group, and the like. Carboxyl



group, acetyl group, carbamoyl group, N,N-dimethylcarbamoyl group, and the like are preferred examples, and carboxyl group is particularly preferred.

Preferred examples of the group Xa include oxo group, thioxo group, fluorine atom, chlorine atom, trifluoromethyl group, methyl group, ethyl group, propyl group, 2-hydroxyethyl group, carboxymethyl group, 2-carboxyethyl group, N,N-dimethylcarbamoylmethyl group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, carboxymethyloxy group, 2-carboxyethyloxy group, N,N-dimethylcarbamoylmethyloxy group, amino group, methylamino group, dimethylamino group, 2-hydroxyethylamino group, carbamoylamino group, acetylamino group, furan-2-carboxyamino group, 2-hydroxyacetylamino group, 2-aminoacetylamino group, methylsulfonylamino group, (N,N-dimethylsulfamoyl)amino group, methanesulfonyl group, sulfamoyl group, N-methylsulfamoyl group, N,N-dimethylsulfamoyl group, carboxyl group, acetyl group, carbamoyl group, N,N-dimethylcarbamoyl group, and the like. Particularly preferred examples of the group Xa include oxo group, methyl group, ethyl group, propyl group, 2-hydroxyethyl group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, amino group, N-methylamino group, N,N-dimethylamino group, 2-hydroxyethylamino group, carboxyl group, and the like. Preferred examples of the group Xa which may substitute on nitrogen atom include methyl group, ethyl group, propyl group, hydroxymethyl group, 2-hydroxyethyl group, carboxymethyl group, 2-carboxyethyl group, and N,N-dimethylcarbamoylmethyl group. Among them, particularly preferred examples are methyl group, ethyl group, propyl group, and 2-hydroxyethyl group.

Preferred examples of AR substituted with the group Xa or unsubstituted AR include naphthalen-1-yl group, naphthalen-2-yl group, 6-fluoronaphthalen-2-yl group, 6-chloronaphthalen-2-yl group, 6-(trifluoromethyl)naphthalen-2-yl group, 5-hydroxynaphthalen-1-yl group, 5-hydroxynaphthalen-2-yl group,

6-hydroxynaphthalen-1-yl group, 6-hydroxynaphthalen-2-yl group,  
 7-hydroxynaphthalen-1-yl group, 7-hydroxynaphthalen-2-yl group,  
 5-methoxynaphthalen-1-yl group, 5-methoxynaphthalen-2-yl group,  
 6-methoxynaphthalen-1-yl group, 6-methoxynaphthalen-2-yl group,  
 7-methoxynaphthalen-1-yl group, 7-methoxynaphthalen-2-yl group,  
 5-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl  
 group, 7-(2-hydroxyethyloxy)naphthalen-2-yl group,  
 5-(carboxymethyloxy)naphthalen-2-yl group, 6-(carboxymethyloxy)naphthalen-2-yl  
 group, 7-(carboxymethyloxy)naphthalen-2-yl group,  
 5-(N,N-dimethylcarbamoylmethyloxy)naphthalen-2-yl group,  
 6-(N,N-dimethylcarbamoylmethyloxy)naphthalen-2-yl group,  
 7-(N,N-dimethylcarbamoylmethyloxy)naphthalen-2-yl group,  
 5-aminonaphthalen-1-yl group, 5-aminonaphthalen-2-yl group,  
 6-aminonaphthalen-1-yl group, 6-aminonaphthalen-2-yl group,  
 7-aminonaphthalen-1-yl group, 7-aminonaphthalen-2-yl group,  
 5-(N-methylamino)naphthalen-1-yl group, 5-(N-methylamino)naphthalen-2-yl group,  
 6-(N-methylamino)naphthalen-1-yl group, 6-(N-methylamino)naphthalen-2-yl group,  
 7-(N-methylamino)naphthalen-1-yl group, 7-(N-methylamino)naphthalen-2-yl group,  
 5-(N,N-dimethylamino)naphthalen-1-yl group,  
 5-(N,N-dimethylamino)naphthalen-2-yl group,  
 6-(N,N-dimethylamino)naphthalen-1-yl group,  
 6-(N,N-dimethylamino)naphthalen-2-yl group,  
 7-(N,N-dimethylamino)naphthalen-1-yl group,  
 7-(N,N-dimethylamino)naphthalen-2-yl group,  
 5-(2-hydroxyethylamino)naphthalen-2-yl group,  
 6-(2-hydroxyethylamino)naphthalen-2-yl group,  
 7-(2-hydroxyethylamino)naphthalen-2-yl group, 5-acetylaminonaphthalen-2-yl

group, 6-acetylamino-naphthalen-2-yl group, 6-(2-aminoacetyl-amino)-naphthalen-2-yl  
 group, 6-(2-hydroxyacetyl-amino)-naphthalen-2-yl group,  
 7-(2-hydroxyacetyl-amino)-naphthalen-2-yl group,  
 6-[(furan-2-carbonyl)-amino]-naphthalen-2-yl group,  
 7-[(furan-2-carbonyl)-amino]-naphthalen-2-yl group,  
 6-[(benzene-2-carbonyl)-amino]-naphthalen-2-yl group,  
 7-[(benzene-2-carbonyl)-amino]-naphthalen-2-yl group,  
 6-carbamoylamino-naphthalen-2-yl group, 6-methylsulfonylamino-naphthalen-2-yl  
 group, 6-sulfamoylamino-naphthalen-2-yl group,  
 6-(N,N-dimethylsulfamoyl-amino)-naphthalen-2-yl group,  
 6-methanesulfonyl-naphthalen-2-yl group, 6-sulfamoyl-naphthalen-2-yl group,  
 6-(N-methylsulfamoyl)-naphthalen-2-yl group,  
 6-(N,N-dimethylsulfamoyl)-naphthalen-2-yl group, 6-carboxy-naphthalen-2-yl group,  
 benzo[b]furan-4-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-4-yl  
 group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-4-yl group,  
 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-4-yl group,  
 2,3-dimethylbenzo[b]furan-5-yl group, 2-carboxybenzo[b]furan-4-yl group,  
 2-carboxybenzo[b]furan-5-yl group, 2-carboxy-3-methylbenzo[b]furan-4-yl group,  
 2-carboxy-3-methylbenzo[b]furan-5-yl group, 3-acetylbenzo[b]furan-4-yl group,  
 3-acetylbenzo[b]furan-5-yl group, 3-acetyl-2-methylbenzo[b]furan-4-yl group,  
 3-acetyl-2-methylbenzo[b]furan-5-yl group, 3-hydroxymethylbenzo[b]furan-4-yl  
 group, 3-hydroxymethylbenzo[b]furan-5-yl group,  
 3-hydroxymethyl-2-methylbenzo[b]furan-4-yl group,  
 3-hydroxymethyl-2-methylbenzo[b]furan-5-yl group, benzo[b]thiophen-4-yl group,  
 benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-4-yl group,  
 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-4-yl group,  
 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-4-yl group,

2,3-dimethylbenzo[b]thiophen-5-yl group, 2-carboxybenzo[b]thiophen-4-yl group,  
 2-carboxybenzo[b]thiophen-5-yl group, 2-carboxy-3-methylbenzo[b]thiophen-4-yl  
 group, 2-carboxy-3-methylbenzo[b]thiophen-5-yl group,  
 3-acetylbenzo[b]thiophen-4-yl group, 3-acetylbenzo[b]thiophen-5-yl group,  
 3-acetyl-2-methylbenzo[b]thiophen-4-yl group,  
 3-acetyl-2-methylbenzo[b]thiophen-5-yl group,  
 3-hydroxymethylbenzo[b]thiophen-4-yl group,  
 3-hydroxymethylbenzo[b]thiophen-5-yl group,  
 3-hydroxymethyl-2-methylbenzo[b]thiophen-4-yl group,  
 3-hydroxymethyl-2-methylbenzo[b]thiophen-5-yl group, 1H-indol-4-yl group,  
 1H-indol-5-yl group, 2-methyl-1H-indol-4-yl group, 2-methyl-1H-indol-5-yl group,  
 3-methyl-1H-indol-4-yl group, 3-methyl-1H-indol-5-yl group,  
 2,3-dimethyl-1H-indol-4-yl group, 2,3-dimethyl-1H-indol-5-yl group,  
 2-carboxy-1H-indol-4-yl group, 2-carboxy-1H-indol-5-yl group,  
 2-carboxy-3-methyl-1H-indol-4-yl group, 2-carboxy-3-methyl-1H-indol-5-yl group,  
 3-acetyl-1H-indol-4-yl group, 3-acetyl-1H-indol-5-yl group,  
 3-acetyl-2-methyl-1H-indol-4-yl group, 3-acetyl-2-methyl-1H-indol-5-yl group,  
 3-hydroxymethyl-1H-indol-4-yl group, 3-hydroxymethyl-1H-indol-5-yl group,  
 3-hydroxymethyl-2-methyl-1H-indol-4-yl group,  
 3-hydroxymethyl-2-methyl-1H-indol-5-yl group, 1-methyl-1H-indol-4-yl group,  
 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-4-yl group,  
 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-4-yl group,  
 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-4-yl group,  
 1,2,3-trimethyl-1H-indol-5-yl group, 2-carboxy-1-methyl-1H-indol-4-yl group,  
 2-carboxy-1-methyl-1H-indol-5-yl group, 2-carboxy-1,3-dimethyl-1H-indol-4-yl group,  
 2-carboxy-1,3-dimethyl-1H-indol-5-yl group, 3-acetyl-1-methyl-1H-indol-4-yl group,  
 3-acetyl-1-methyl-1H-indol-5-yl group, 3-acetyl-1,2-dimethyl-1H-indol-4-yl group,

3-acetyl-1,2-dimethyl-1H-indol-5-yl group, 3-hydroxymethyl-1-methyl-1H-indol-4-yl  
 group, 3-hydroxymethyl-1-methyl-1H-indol-5-yl group,  
 3-hydroxymethyl-1,2-dimethyl-1H-indol-4-yl group,  
 3-hydroxymethyl-1,2-dimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-4-yl group,  
 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-4-yl group,  
 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-4-yl group,  
 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-4-yl group,  
 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 2-carboxy-1-ethyl-1H-indol-4-yl group,  
 2-carboxy-1-ethyl-1H-indol-5-yl group, 2-carboxy-1-ethyl-3-methyl-1H-indol-4-yl  
 group, 2-carboxy-1-ethyl-3-methyl-1H-indol-5-yl group,  
 3-acetyl-1-ethyl-1H-indol-4-yl group, 3-acetyl-1-ethyl-1H-indol-5-yl group,  
 3-acetyl-1-ethyl-2-methyl-1H-indol-4-yl group,  
 3-acetyl-1-ethyl-2-methyl-1H-indol-5-yl group,  
 1-ethyl-3-hydroxymethyl-1H-indol-4-yl group,  
 1-ethyl-3-hydroxymethyl-1H-indol-5-yl group,  
 1-ethyl-3-hydroxymethyl-2-methyl-1H-indol-4-yl group,  
 1-ethyl-3-hydroxymethyl-2-methyl-1H-indol-5-yl group, 1-propyl-1H-indol-4-yl  
 group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-4-yl group,  
 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-4-yl group,  
 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-4-yl group,  
 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 2-carboxy-1-propyl-1H-indol-4-yl group,  
 2-carboxy-1-propyl-1H-indol-5-yl group, 2-carboxy-3-methyl-1-propyl-1H-indol-4-yl  
 group, 2-carboxy-3-methyl-1-propyl-1H-indol-5-yl group,  
 3-acetyl-1-propyl-1H-indol-4-yl group, 3-acetyl-1-propyl-1H-indol-5-yl group,  
 3-acetyl-2-methyl-1-propyl-1H-indol-4-yl group,  
 3-acetyl-2-methyl-1-propyl-1H-indol-5-yl group,  
 3-hydroxymethyl-1-propyl-1H-indol-4-yl group,

3-hydroxymethyl-1-propyl-1H-indol-5-yl group,  
 3-hydroxymethyl-2-methyl-1-propyl-1H-indol-4-yl group,  
 3-hydroxymethyl-2-methyl-1-propyl-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-1H-indol-4-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-2-methyl-1H-indol-4-yl group,  
 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-3-methyl-1H-indol-4-yl group,  
 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group,  
 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-4-yl group,  
 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group,  
 2-carboxy-1-(2-hydroxyethyl)-1H-indol-4-yl group,  
 2-carboxy-1-(2-hydroxyethyl)-1H-indol-5-yl group,  
 2-carboxy-1-(2-hydroxyethyl)-3-methyl-1H-indol-4-yl group,  
 2-carboxy-1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group,  
 3-acetyl-1-(2-hydroxyethyl)-1H-indol-4-yl group,  
 3-acetyl-1-(2-hydroxyethyl)-1H-indol-5-yl group,  
 3-acetyl-1-(2-hydroxyethyl)-2-methyl-1H-indol-4-yl group,  
 3-acetyl-1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-3-hydroxymethyl-1H-indol-4-yl group,  
 1-(2-hydroxyethyl)-3-hydroxymethyl-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-3-hydroxymethyl-2-methyl-1H-indol-4-yl group,  
 1-(2-hydroxyethyl)-3-hydroxymethyl-2-methyl-1H-indol-5-yl group,  
 1-carboxymethyl-1H-indol-4-yl group, 1-carboxymethyl-1H-indol-5-yl group,  
 1-carboxymethyl-2-methyl-1H-indol-4-yl group,  
 1-carboxymethyl-2-methyl-1H-indol-5-yl group,  
 1-carboxymethyl-3-methyl-1H-indol-4-yl group,  
 1-carboxymethyl-3-methyl-1H-indol-5-yl group,

1-carboxymethyl-2,3-dimethyl-1H-indol-4-yl group,  
 1-carboxymethyl-2,3-dimethyl-1H-indol-5-yl group,  
 2-carboxy-1-carboxymethyl-1H-indol-4-yl group,  
 2-carboxy-1-carboxymethyl-1H-indol-5-yl group,  
 2-carboxy-1-carboxymethyl-3-methyl-1H-indol-4-yl group,  
 2-carboxy-1-carboxymethyl-3-methyl-1H-indol-5-yl group,  
 3-acetyl-1-carboxymethyl-1H-indol-4-yl group,  
 3-acetyl-1-carboxymethyl-1H-indol-5-yl group,  
 3-acetyl-1-carboxymethyl-2-methyl-1H-indol-4-yl group,  
 3-acetyl-1-carboxymethyl-2-methyl-1H-indol-5-yl group,  
 1-carboxymethyl-3-hydroxymethyl-1H-indol-4-yl group,  
 1-carboxymethyl-3-hydroxymethyl-1H-indol-5-yl group,  
 1-carboxymethyl-3-hydroxymethyl-2-methyl-1H-indol-4-yl group,  
 1-carboxymethyl-3-hydroxymethyl-2-methyl-1H-indol-5-yl group, benzothiazol-6-yl  
 group, 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group,  
 2-aminobenzothiazol-6-yl group, 2-(N-methylamino)benzothiazol-6-yl group,  
 2-(N,N-dimethylamino)benzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl  
 group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group,  
 2-methylquinolin-3-yl group, quinolin-6-yl group, 2-methylquinolin-6-yl group,  
 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group,  
 3-methylbenzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group,  
 3-methyl-1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group,  
 1,3-dimethyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group,  
 1-ethyl-3-methyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group,  
 3-methyl-1-propyl-1H-indazol-5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group,

1-(2-hydroxyethyl)-3-methyl-1H-indazol-5-yl group,  
 1-(carboxymethyl)-1H-indazol-5-yl group,  
 1-(carboxymethyl)-3-methyl-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group,  
 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group,  
 benzo[c]isothiazol-5-yl group, 3-methylbenzo[c]isothiazol-5-yl group,  
 2-methyl-2H-indazol-5-yl group, 2,3-dimethyl-2H-indazol-5-yl group,  
 2-ethyl-2H-indazol-5-yl group, 2-ethyl-3-methyl-2H-indazol-5-yl group,  
 2-propyl-2H-indazol-5-yl group, 3-methyl-2-propyl-2H-indazol-5-yl group,  
 2-(2-hydroxyethyl)-2H-indazol-5-yl group,  
 2-(2-hydroxyethyl)-3-methyl-2H-indazol-5-yl group,  
 2-(carboxymethyl)-2H-indazol-5-yl group,  
 2-(carboxymethyl)-3-methyl-2H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group,  
 2-methyl-imidazo[1,2-a]pyridin-6-yl group, 3-methyl-imidazo[1,2-a]pyridin-6-yl  
 group, 2,3-dimethyl-imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl  
 group, 2-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 3-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1,2-dimethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1,3-dimethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 2,3-dimethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1,2,3-trimethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-ethyl-2-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-ethyl-3-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-ethyl-2,3-dimethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 2-methyl-1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,



3-methyl-1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 2,3-dimethyl-1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-(2-hydroxyethyl)-2-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-(2-hydroxyethyl)-3-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 2,3-dimethyl-1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-(carboxymethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-(carboxymethyl)-2-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-(carboxymethyl)-3-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-(carboxymethyl)-2,3-dimethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 isoquinolin-6-yl group, 1-methylisoquinolin-6-yl group,  
 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, cinnolin-5-yl group,  
 quinazolin-6-yl group, quinazolin-7-yl group, quinazolin-5-yl group,  
 2-methylquinazolin-6-yl group, quinoxalin-6-yl group, quinoxalin-5-yl group,  
 2-methylquinoxalin-6-yl group, 1H-benzimidazol-5-yl group, 1H-benzimidazol-4-yl  
 group, 1-methyl-1H-benzimidazol-5-yl group, 2-methyl-1H-benzimidazol-5-yl group,  
 1,2-dimethyl-1H-benzimidazol-5-yl group, benzoxazol-5-yl group, benzoxazol-6-yl  
 group, benzoxazol-4-yl group, benzoxazol-7-yl group, 2-methylbenzoxazol-5-yl group,  
 1H-pyrrolo[3,2-b]pyridin-5-yl group, 1H-pyrrolo[3,2-b]pyridin-6-yl group,  
 1-methyl-1H-pyrrolo[3,2-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[3,2-b]pyridin-5-yl  
 group, 2-methyl-1H-pyrrolo[3,2-b]pyridin-5-yl group,  
 3-methyl-1H-pyrrolo[3,2-b]pyridin-5-yl group,  
 1,3-dimethyl-1H-pyrrolo[3,2-b]pyridin-5-yl group, benzo[1,2,5]thiadiazol-5-yl group,  
 benzo[1,2,5]thiadiazol-4-yl group, 1H-benzotriazol-5-yl group, 1H-benzotriazol-4-yl  
 group, 1-methyl-1H-benzotriazol-5-yl group, 1-ethyl-1H-benzotriazol-5-yl group,  
 1,3-dihydropyrrolo[2,3-b]pyridin-2-on-5-yl group,  
 1,3-dihydropyrrolo[2,3-b]pyridin-2-on-4-yl group,

1-methyl-1,3-dihydropyrrolo[2,3-b]pyridin-2-on-5-yl group,  
 1,3-dihydrobenzimidazol-2-on-5-yl group, 1,3-dihydrobenzimidazol-2-on-4-yl group,  
 1-methyl-1,3-dihydrobenzimidazol-2-on-5-yl group,  
 1,3-dihydrobenzimidazole-2-thion-5-yl group, 1,3-dihydrobenzimidazole-2-thion-4-yl  
 group, 1-methyl-1,3-dihydrobenzimidazole-2-thion-5-yl group,  
 3H-benzoxazol-2-on-6-yl group, 3H-benzoxazol-2-on-7-yl group,  
 3H-benzoxazol-2-on-5-yl group, 3H-benzoxazol-2-on-4-yl group,  
 3-methyl-3H-benzoxazol-2-on-6-yl group, 3H-benzoxazole-2-thion-6-yl group,  
 3H-benzoxazole-2-thion-7-yl group, 3H-benzoxazole-2-thion-5-yl group,  
 3H-benzoxazole-2-thion-4-yl group, 3-methyl-3H-benzoxazole-2-thion-6-yl group,  
 phthalazin-6-yl group, phthalazin-5-yl group, [1,8]naphthalidin-3-yl group,  
 [1,8]naphthalidin-4-yl group, [1,5]naphthalidin-3-yl group, [1,5]naphthalidin-4-yl  
 group, 1H-pyrrolo[3,2-c]pyridin-6-yl group, 1H-pyrrolo[3,2-c]pyridin-4-yl group,  
 1-methyl-1H-pyrrolo[3,2-c]pyridin-6-yl group, 1-ethyl-1H-pyrrolo[3,2-c]pyridin-6-yl  
 group, 2-methyl-1H-pyrrolo[3,2-c]pyridin-6-yl group,  
 3-methyl-1H-pyrrolo[3,2-c]pyridin-6-yl group,  
 1,3-dimethyl-1H-pyrrolo[3,2-c]pyridin-6-yl group, 1H-pyrrolo[2,3-c]pyridin-5-yl  
 group, 1H-pyrrolo[2,3-c]pyridin-4-yl group, 1-methyl-1H-pyrrolo[2,3-c]pyridin-5-yl  
 group, 1-ethyl-1H-pyrrolo[2,3-c]pyridin-5-yl group,  
 2-methyl-1H-pyrrolo[2,3-c]pyridin-5-yl group,  
 3-methyl-1H-pyrrolo[2,3-c]pyridin-5-yl group,  
 1,3-dimethyl-1H-pyrrolo[2,3-c]pyridin-5-yl group, 1H-pyrazolo[4,3-b]pyridin-5-yl  
 group, 1H-pyrazolo[4,3-b]pyridin-6-yl group,  
 1-methyl-1H-pyrazolo[4,3-b]pyridin-5-yl group,  
 1-ethyl-1H-pyrazolo[4,3-b]pyridin-5-yl group,  
 3-methyl-1H-pyrazolo[4,3-b]pyridin-5-yl group,  
 1,3-dimethyl-1H-pyrazolo[4,3-b]pyridin-5-yl group, 1H-pyrazolo[4,3-c]pyridin-6-yl

group, 1H-pyrazolo[4,3-c]pyridin-4-yl group,  
 1-methyl-1H-pyrazolo[4,3-c]pyridin-6-yl group,  
 1-ethyl-1H-pyrazolo[4,3-c]pyridin-6-yl group,  
 3-methyl-1H-pyrazolo[4,3-c]pyridin-6-yl group,  
 1,3-dimethyl-1H-pyrazolo[4,3-c]pyridin-6-yl group, 1H-pyrazolo[3,4-c]pyridin-5-yl  
 group, 1H-pyrazolo[3,4-c]pyridin-4-yl group,  
 1-methyl-1H-pyrazolo[3,4-c]pyridin-5-yl group,  
 1-ethyl-1H-pyrazolo[3,4-c]pyridin-5-yl group,  
 3-methyl-1H-pyrazolo[3,4-c]pyridin-5-yl group,  
 1,3-dimethyl-1H-pyrazolo[3,4-c]pyridin-5-yl group, 1H-pyrazolo[3,4-b]pyridin-5-yl  
 group, 1H-pyrazolo[3,4-b]pyridin-4-yl group,  
 1-methyl-1H-pyrazolo[3,4-b]pyridin-5-yl group,  
 1-ethyl-1H-pyrazolo[3,4-b]pyridin-5-yl group,  
 3-methyl-1H-pyrazolo[3,4-b]pyridin-5-yl group,  
 1,3-dimethyl-1H-pyrazolo[3,4-b]pyridin-5-yl group, [1,2,4]triazolo[4,3-a]pyridin-6-yl  
 group, [1,2,4]triazolo[4,3-a]pyridin-7-yl group,  
 3-methyl[1,2,4]triazolo[4,3-a]pyridin-6-yl group, thieno[3,2-c]pyridin-2-yl group,  
 thieno[3,2-c]pyridin-3-yl group, thieno[3,2-c]pyridin-6-yl group,  
 2-methylthieno[3,2-c]pyridin-2-yl group, 3-methylthieno[3,2-c]pyridin-2-yl group,  
 thieno[3,2-b]pyridin-2-yl group, thieno[3,2-b]pyridin-3-yl group,  
 thieno[3,2-b]pyridin-5-yl group, thieno[3,2-b]pyridin-6-yl group,  
 2-methylthieno[3,2-b]pyridin-2-yl group, 3-methylthieno[3,2-b]pyridin-2-yl group,  
 1H-thieno[3,2-c]pyrazol-5-yl group, 1H-thieno[3,2-c]pyrazol-4-yl group,  
 1-methyl-1H-thieno[3,2-c]pyrazol-5-yl group, 1-ethyl-1H-thieno[3,2-c]pyrazol-5-yl  
 group, 3-methyl-1H-thieno[3,2-c]pyrazol-5-yl group,  
 1,3-dimethyl-1H-thieno[3,2-c]pyrazol-5-yl group, benzo[d]isoxazol-5-yl group,  
 benzo[d]isoxazol-4-yl group, benzo[d]isoxazol-6-yl group, benzo[d]isoxazol-7-yl group,

3-methylbenzo[d]isoxazol-5-yl group, benzo[c]isoxazol-5-yl group, benzo[c]isoxazol-4-yl group, benzo[c]isoxazol-6-yl group, benzo[c]isoxazol-7-yl group, 3-methylbenzo[c]isoxazol-5-yl group, indolizin-7-yl group, indolizin-6-yl group, indolizine-8-yl group, 1,3-dihydroindol-2-on-5-yl group, 1,3-dihydroindol-2-on-4-yl group, 1,3-dihydroindol-2-on-6-yl group, 1-methyl-1,3-dihydro-indol-2-on-5-yl group, 1H-pyrazolo[3,4-d]thiazol-5-yl group, 2H-isoindol-5-yl group, 2H-isoindol-4-yl group, 2-methyl-2H-isoindol-5-yl group, 4H-chromen-6-yl group, 4H-chromen-5-yl group, chromen-4-on-7-yl group, chromen-4-on-6-yl group, and the like.

Particularly preferred examples include naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group,

2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group, 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, benzoxazol-5-yl group, and the like.

Particularly preferred examples include naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, benzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group,

imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, benzoxazol-5-yl group, and the like.

In the formula (I), the group Y is defined to be hydrogen atom, a lower alkyl group having 1 to 4 carbon atoms,  $-(CH_2)_mN(R^{18})(R^{19})$ , or  $-C(R^{20})_2OC(O)A^3R^{21}$ , and among them, hydrogen atom is particularly preferred.

Examples of the lower alkyl group having 1 to 4 carbon atoms include methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, and the like. Among them, methyl group, and ethyl group are particularly preferred.

Symbol m in  $-(CH_2)_mN(R^{18})(R^{19})$  is defined to be an integer of 2 or 3.  $R^{18}$  is the same as  $R^{19}$ , or binds to  $R^{19}$  to represent a saturated nitrogen-containing cycloalkyl group forming a 3- to 6-membered ring together with nitrogen atom, or form morpholino group together with nitrogen atom, and  $R^{19}$  is defined to be methyl group, ethyl group, or propyl group. Examples of  $-(CH_2)_mN(R^{18})(R^{19})$  include 2-(N,N-dimethylamino)ethyl group, 2-(N,N-diethylamino)ethyl group, 2-(N,N-dipropylamino)ethyl group, 3-(N,N-dimethylamino)propyl group, 3-(N,N-diethylamino)propyl group, 2-(N,N-dipropylamino)propyl group, 2-pyrrolidin-1-ylethyl group, 2-piperidin-1-ylethyl group, 2-morpholin-4-ylethyl group, 3-pyrrolidin-1-ylpropyl group, 3-piperidin-1-ylpropyl group, 3-morpholin-4-ylpropyl group, and the like.

$R^{20}$  in  $-C(R^{20})_2OC(O)A^3R^{21}$  is defined to be hydrogen atom, methyl group, ethyl group, or propyl group.  $R^{21}$  is defined to be a lower alkyl group having 1 to 4 carbon atoms, a cyclic saturated alkyl group having 3 to 6 carbon atoms group, or phenyl group. Examples of the lower alkyl group having 1 to 4 carbon atoms include methyl group, ethyl group, propyl group, isopropyl group, butyl group,

isobutyl group, t-butyl group, and the like, and examples of the cyclic saturated alkyl group having 3 to 6 carbon atoms group include cyclopropyl group, cyclobutyl group, cyclopentyl group, and cyclohexyl group.  $A^3$  is defined to be a single bond, or oxygen atom. Examples of  $-C(R^{20})_2OC(O)A^3R^{21}$  include acetoxymethyl group, propionyloxymethyl group, butyryloxymethyl group, (2-methylpropionyl)oxymethyl group, (2,2-dimethylpropionyl)oxymethyl group, cyclopropionyloxymethyl group, cyclopentanoyloxymethyl group, cyclohexanoyloxymethyl group, phenylcarboxymethyl group, 1-acetoxy-1-methylethyl group, 1-methyl-1-(2-methylpropionyloxy)ethyl group, 1-cyclopentanoyloxy-1-methylethyl group, 1-cyclohexanoyloxy-1-methylethyl group, methoxycarbonyloxymethyl group, ethoxycarbonyloxymethyl group, isopropylloxycarbonyloxymethyl group, t-butyloxycarbonyloxymethyl group, cyclopropylloxycarbonyloxymethyl group, cyclopentylloxycarbonyloxymethyl group, cyclohexylloxycarbonyloxymethyl group, phenylloxycarbonyloxymethyl group, 1-methoxycarbonyloxy-1-methylethyl group, 1-ethoxycarbonyloxy-1-methylethyl group, 1-isopropylloxycarbonyloxy-1-methylethyl group, 1-t-butyloxycarbonyloxy-1-methylethyl group, 1-cyclopropylloxycarbonyloxy-1-methylethyl group, 1-cyclopentylloxycarbonyloxy-1-methylethyl group, 1-cyclohexylloxycarbonyloxy-1-methylethyl group, 1-methyl-1-phenylloxycarbonyloxyethyl group, and the like.

In a preferred embodiment of the present invention, the compound represented by the formula (I) or a salt thereof satisfies all of the following requirements.

Link represents  $-(CH_2)_n-$ , symbol n represents an integer of 1 to 3.

AR binds to  $C^2$ , Rs binds to any of the atoms  $C^3$ ,  $C^4$  and  $C^5$ , and a ring-constituting carbon atom to which Rs does not bind among  $C^3$ ,  $C^4$ , and  $C^5$  may be replaced with V.

V represents nitrogen atom, or carbon atom substituted with Zx, and Zx represents a group as any one of fluorine atom, chlorine atom, bromine atom, nitro group, methyl group, hydroxyl group, methoxy group, amino group, N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, and N,N-dimethylsulfamoylamino group.

Rs represents -D-Rx, or -N(Ry)(Rz). D represents oxygen atom, or sulfur atom. Rx represents butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-cyclopentylethyl group, or 2-cyclohexylethyl group, or represents Rb or Rc. Q in Rb represents a group as any one of phenyl group, thienyl group, furyl group, pyridyl group, oxazolyl group, naphthyl group, tetrahydronaphthyl group, indanyl group, indolyl group, and dihydrobenzodioxyl group. A<sup>2</sup> represents a single bond, oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)- (provided that when A<sup>2</sup> represents oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)-, A<sup>1</sup> represents ethylene). R<sup>2</sup> and R<sup>3</sup> independently represent hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, dimethylamino group, acetylamino group, or methylsulfonylamino group (provided that when Q represents phenyl group, A<sup>1</sup> represents a single bond, or unsubstituted methylene, and A<sup>2</sup> represents a single bond, one of R<sup>2</sup> and R<sup>3</sup> represents a substituent other than hydrogen atom). Symbol p in Rc represents an integer of 2 or 3, and A<sup>4</sup> represents a single bond or methylene. A<sup>5</sup> represents -C(O)-, -C(S)-, or -S(O)<sub>2</sub>-. Rd represents hydrogen atom, or a group as any one of methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group, cyclopropylmethyl group, cyclopentyl group, cyclopentylmethyl group, cyclohexyl group, cyclohexylmethyl group, phenyl group,



4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, and pyridin-4-yl group. Re represents a group as any one of methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, phenylmethyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, furan-2-yl group, furan-3-yl group, thiophen-2-yl group, thiophen-3-yl group, methoxy group, ethoxy group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenyloxy group, 4-methylphenyloxy group, 4-chlorophenyloxy group, 4-fluorophenyloxy group, thiomethoxy group, amino group, N-methylamino group, N,N-dimethylamino group, N-ethylamino group, N,N-diethylamino group, N-propylamino group, N-isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group, N-(4-fluorophenyl)amino group, N-(pyridin-2-yl)amino group, N-(pyridin-3-yl)amino group, N-(pyridin-4-yl)amino group, N-(furan-2-yl)amino group, N-(furan-3-yl)amino group, N-(thiophen-2-yl)amino group, N-(thiophen-3-yl)amino group, pyrrolidino group, piperidino group, morpholino group, methyloxycarbonylamino group, and ethyloxycarbonylamino group.

Rz represents a group as any one of butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl

group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group,  
 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl  
 group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group,  
 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group,  
 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group,  
 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group,  
 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group,  
 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl  
 group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl  
 group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group,  
 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group,  
 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl  
 group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group,  
 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group,  
 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl  
 group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group,  
 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group,  
 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group,  
 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group,  
 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group,  
 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group,  
 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group,  
 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group,  
 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group,  
 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group,  
 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group,  
 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group,

2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group,  
 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group,  
 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group,  
 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl  
 group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group,  
 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group,  
 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group,  
 2-(N-ethyl-N-phenylamino)ethyl group, isobutyryl group, isopropylthiocarbonyl  
 group, isopropylsulfonyl group, valeryl group, butylthiocarbonyl group, isovaleryl  
 group, isobutylthiocarbonyl group, pivaloyl group, t-butylthiocarbonyl group,  
 cyclopropylcarbonyl group, cyclopropylthiocarbonyl group, cyclopentylcarbonyl  
 group, cyclopentylthiocarbonyl group, cyclohexylcarbonyl group,  
 cyclohexylthiocarbonyl group, cyclopentylmethylcarbonyl group,  
 cyclopentylmethylthiocarbonyl group, cyclohexylmethylcarbonyl group,  
 cyclohexylmethylthiocarbonyl group, benzoyl group, thiobenzoyl group,  
 phenylsulfonyl group, 4-methylphenylcarbonyl group, 4-methylphenylthiocarbonyl  
 group, 4-methylphenylsulfonyl group, 4-chlorophenylcarbonyl group,  
 4-chlorophenylthiocarbonyl group, 4-fluorophenylcarbonyl group,  
 4-fluorophenylthiocarbonyl group, isopropoxy carbonyl group,  
 N-isopropylcarbamoyl group, N-isopropylthiocarbamoyl group, butyloxycarbonyl  
 group, N-butylcarbamoyl group, N-butylthiocarbamoyl group, isobutyloxycarbonyl  
 group, N-isobutylcarbamoyl group, N-isobutylthiocarbamoyl group,  
 t-butyloxycarbonyl group, N-t-butylcarbamoyl group, N-t-butylthiocarbamoyl group,  
 cyclopropyloxycarbonyl group, N-cyclopropylcarbamoyl group,  
 N-cyclopropylthiocarbamoyl group, cyclopentyloxycarbonyl group,  
 N-cyclopentylcarbamoyl group, N-cyclopentylthiocarbamoyl group,  
 cyclohexyloxycarbonyl group, N-cyclohexylcarbamoyl group,

N-cyclohexylthiocarbamoyl group, cyclopentylmethyloxycarbonyl group, cyclohexylmethyloxycarbonyl group, phenyloxycarbonyl group, N-phenylcarbamoyl group, N-phenylthiocarbamoyl group, 4-methylphenyloxycarbonyl group, N-(4-methylphenyl)carbamoyl group, N-(4-methylphenyl)thiocarbamoyl group, 4-chlorophenyloxycarbonyl group, N-(4-chlorophenyl)carbamoyl group, N-(4-chlorophenyl)thiocarbamoyl group, 4-fluorophenyloxycarbonyl group, N-(4-fluorophenyl)carbamoyl group, N-(4-fluorophenyl)thiocarbamoyl group, (pyrrolidino-1-yl)carbonyl group, (piperidino-1-yl)carbonyl group, and (morpholino-4-yl)carbonyl group. Ry represents hydrogen atom, methyl group, ethyl group, or isobutyl group, or binds to Rz to form pyrrolidino group, piperidino group, piperazino group, morpholino group, pyrrol-1-yl group, imidazol-1-yl group, or pyrazol-1-yl group together with nitrogen atom.

AR represents naphthalen-2-yl group, naphthalen-1-yl group, benzofuran-5-yl group, benzofuran-4-yl group, benzofuran-2-yl group, benzo[b]thiophen-5-yl group, benzo[b]thiophen-4-yl group, benzo[b]thiophen-2-yl group, indol-5-yl group, indol-4-yl group, indol-6-yl group, benzothiazol-6-yl group, benzothiazol-7-yl group, benzothiazol-5-yl group, benzothiazol-4-yl group, dihydro-3H-benzothiazol-6-yl group, dihydro-3H-benzothiazol-7-yl group, dihydro-3H-benzothiazol-5-yl group, dihydro-3H-benzothiazol-4-yl group, quinolin-6-yl group, quinolin-3-yl group, quinolin-5-yl group, quinolin-7-yl group, dihydro-1H-quinolin-6-yl group, dihydro-1H-quinolin-5-yl group, benzo[d]isothiazol-5-yl group, benzo[d]isothiazol-4-yl group, benzo[d]isothiazol-6-yl group, benzo[d]isothiazol-7-yl group, 1H-indazol-5-yl group, 1H-indazol-4-yl group, 1H-indazol-6-yl group, benzo[c]isothiazol-5-yl group, benzo[c]isothiazol-4-yl group, benzo[c]isothiazol-6-yl group, benzo[c]isothiazol-7-yl group, 2H-indazol-5-yl group, 2H-indazol-4-yl group, 2H-indazol-6-yl group, imidazo[1,2-a]pyridin-6-yl group, imidazo[1,2-a]pyridin-7-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group,

1H-pyrrolo[2,3-b]pyridin-4-yl group, isoquinolin-6-yl group, isoquinolin-3-yl group,  
 isoquinolin-5-yl group, isoquinolin-7-yl group, dihydro-2H-isoquinolin-6-yl group,  
 dihydro-2H-isoquinolin-5-yl group, cinnolin-6-yl group, cinnolin-5-yl group,  
 quinazolin-6-yl group, quinazolin-7-yl group, quinazolin-5-yl group, quinoxalin-2-yl  
 group, quinoxalin-6-yl group, quinoxalin-5-yl group, 1H-benzimidazol-5-yl group,  
 1H-benzimidazol-4-yl group, benzoxazol-5-yl group, benzoxazol-6-yl group,  
 benzoxazol-4-yl group, benzoxazol-7-yl group, 1H-pyrrolo[3,2-b]pyridin-5-yl group,  
 1H-pyrrolo[3,2-b]pyridin-6-yl group, benzo[1,2,5]thiadiazol-5-yl group,  
 benzo[1,2,5]thiadiazol-4-yl group, 1H-benzotriazol-5-yl group, 1H-benzotriazol-4-yl  
 group, 1,3-dihydropyrrolo[2,3-b]pyridin-5-yl group,  
 1,3-dihydropyrrolo[2,3-b]pyridin-4-yl group, 1,3-dihydrobenzimidazol-5-yl group,  
 1,3-dihydrobenzimidazol-4-yl group, dihydro-3H-benzoxazol-6-yl group,  
 dihydro-3H-benzoxazol-7-yl group, dihydro-3H-benzoxazol-5-yl group,  
 dihydro-3H-benzoxazol-4-yl group, phthalazin-6-yl group, phthalazin-5-yl group,  
 [1,8]naphthalidin-3-yl group, [1,8]naphthalidin-4-yl group, [1,5]naphthalidin-3-yl  
 group, [1,5]naphthalidin-4-yl group, 1H-pyrrolo[3,2-c]pyridin-6-yl group,  
 1H-pyrrolo[3,2-c]pyridin-4-yl group, 1H-pyrrolo[2,3-c]pyridin-5-yl group,  
 1H-pyrrolo[2,3-c]pyridin-4-yl group, 1H-pyrazolo[4,3-b]pyridin-5-yl group,  
 1H-pyrazolo[4,3-b]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-6-yl group,  
 1H-pyrazolo[4,3-c]pyridin-4-yl group, 1H-pyrazolo[3,4-c]pyridin-5-yl group,  
 1H-pyrazolo[3,4-c]pyridin-4-yl group, 1H-pyrazolo[3,4-b]pyridin-5-yl group,  
 1H-pyrazolo[3,4-b]pyridin-4-yl group, [1,2,4]triazolo[4,3-a]pyridin-6-yl group,  
 [1,2,4]triazolo[4,3-a]pyridin-7-yl group, thieno[3,2-c]pyridin-2-yl group,  
 thieno[3,2-c]pyridin-3-yl group, thieno[3,2-c]pyridin-6-yl group,  
 thieno[3,2-b]pyridin-2-yl group, thieno[3,2-b]pyridin-3-yl group,  
 thieno[3,2-b]pyridin-5-yl group, thieno[3,2-b]pyridin-6-yl group,  
 1H-thieno[3,2-c]pyrazol-5-yl group, 1H-thieno[3,2-c]pyrazol-4-yl group,

benzo[d]isoxazol-5-yl group, benzo[d]isoxazol-4-yl group, benzo[d]isoxazol-6-yl group, benzo[d]isoxazol-7-yl group, benzo[c]isoxazol-5-yl group, benzo[c]isoxazol-4-yl group, benzo[c]isoxazol-6-yl group, benzo[c]isoxazol-7-yl group, indolizin-7-yl group, indolizin-6-yl group, indolizine-8-yl group, 1,3-dihydroindol-5-yl group, 1,3-dihydroindol-4-yl group, 1,3-dihydroindol-6-yl group, 1H-pyrazolo[3,4-d]thiazol-5-yl group, 2H-isoindol-5-yl group, 2H-isoindol-4-yl group, [1,2,4]triazolo[1,5-a]pyrimidin-6-yl group, 1H-pyrazolo[3,4-b]pyrazin-5-yl group, 1H-imidazo[4,5-b]pyrazin-5-yl group, 7H-purin-2-yl group, 4H-chromen-6-yl group, or 4H-chromen-5-yl group (these groups may be substituted with one of Xa or two or more of the same or different Xa). The substituent Xa represents a group as any one of oxo group, thioxo group, fluorine atom, chlorine atom, trifluoromethyl group, methyl group, ethyl group, propyl group, 2-hydroxyethyl group, carboxymethyl group, 2-carboxyethyl group, N,N-dimethylcarbamoylmethyl group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, carboxymethyloxy group, 2-carboxyethyloxy group, N,N-dimethylcarbamoylmethyloxy group, amino group, methylamino group, dimethylamino group, 2-hydroxyethylamino group, carbamoylamino group, acetylamino group, furan-2-carboxyamino group, 2-hydroxyacetylamino group, 2-aminoacetylamino group, methylsulfonylamino group, (N,N-dimethylsulfamoyl)amino group, methanesulfonyl group, sulfamoyl group, N-methylsulfamoyl group, N,N-dimethylsulfamoyl group, carboxyl group, acetyl group, carbamoyl group, and N,N-dimethylcarbamoyl group.

The group Y represents hydrogen atom, methyl group, or ethyl group.

In another preferred embodiment of the present invention, the compound represented by the formula (I) or a salt thereof satisfies all of the following requirements.

Link represents  $-(CH_2)_n-$ , symbol n represents an integer of 1 to 3.

AR binds to C<sup>3</sup>, Rs binds to any of the atoms C<sup>4</sup>, C<sup>5</sup>, and C<sup>6</sup>, and a

ring-constituting carbon atom to which Rs does not bind among C<sup>4</sup>, C<sup>5</sup>, and C<sup>6</sup> may be replaced with V.

V represents nitrogen atom, or carbon atom substituted with Zx, and Zx represents a group as any one of fluorine atom, chlorine atom, bromine atom, nitro group, methyl group, hydroxyl group, methoxy group, amino group, N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, and N,N-dimethylsulfamoylamino group.

Rs represents -D-Rx, or -N(Ry)(Rz). D represents oxygen atom, or sulfur atom. Rx represents butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-cyclopentylethyl group, or 2-cyclohexylethyl group, or represents Rb, or Rc. Q in Rb represents a group as any one of phenyl group, thienyl group, furyl group, pyridyl group, oxazolyl group, naphthyl group, tetrahydronaphthyl group, indanyl group, indolyl group, and dihydrobenzodioxyl group. A<sup>2</sup> represents a single bond, oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)- (provided that when A<sup>2</sup> represents oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)-, A<sup>1</sup> represents ethylene). R<sup>2</sup> and R<sup>3</sup> independently represent hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, dimethylamino group, acetylamino group, or methylsulfonylamino group (provided that when Q represents phenyl group, A<sup>1</sup> represents a single bond, or unsubstituted methylene, and A<sup>2</sup> represents a single bond, one of R<sup>2</sup> and R<sup>3</sup> represents a substituent other than hydrogen atom). Symbol p in Rc represents an integer of 2 or 3, and A<sup>4</sup> represents a single bond or methylene. A<sup>5</sup> represents -C(O)-, -C(S)-, or -S(O)<sub>2</sub>-. Rd represents hydrogen atom, or a group as any one of methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl

group, cyclopropyl group, cyclopropylmethyl group, cyclopentyl group, cyclopentylmethyl group, cyclohexyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, and pyridin-4-yl group. Re represents a group as any one of methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, phenylmethyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, furan-2-yl group, furan-3-yl group, thiophen-2-yl group, thiophen-3-yl group, methoxy group, ethoxy group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenoxy group, 4-methylphenoxy group, 4-chlorophenoxy group, 4-fluorophenoxy group, thiomethoxy group, amino group, N-methylamino group, N,N-dimethylamino group, N-ethylamino group, N,N-diethylamino group, N-propylamino group, N-isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group, N-(4-fluorophenyl)amino group, N-(pyridin-2-yl)amino group, N-(pyridin-3-yl)amino group, N-(pyridin-4-yl)amino group, N-(furan-2-yl)amino group, N-(furan-3-yl)amino group, N-(thiophen-2-yl)amino group, N-(thiophen-3-yl)amino group, pyrrolidino group, piperidino group, morpholino group, methoxycarbonylamino group, and ethoxycarbonylamino group. Rz represents a group as any of butyl group,



isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group,

2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group,  
 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,  
 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,  
 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group,  
 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group,  
 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group,  
 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group,  
 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group,  
 2-(N-phenyl-N-methylamino)ethyl group, 2-(N-ethyl-N-phenylamino)ethyl group,  
 isobutyryl group, isopropylthiocarbonyl group, isopropylsulfonyl group, valeryl  
 group, butylthiocarbonyl group, isovaleryl group, isobutylthiocarbonyl group,  
 pivaloyl group, t-butylthiocarbonyl group, cyclopropylcarbonyl group,  
 cyclopropylthiocarbonyl group, cyclopentylcarbonyl group, cyclopentylthiocarbonyl  
 group, cyclohexylcarbonyl group, cyclohexylthiocarbonyl group,  
 cyclopentylmethylcarbonyl group, cyclopentylmethylthiocarbonyl group,  
 cyclohexylmethylcarbonyl group, cyclohexylmethylthiocarbonyl group, benzoyl  
 group, thiobenzoyl group, phenylsulfonyl group, 4-methylphenylcarbonyl group,  
 4-methylphenylthiocarbonyl group, 4-methylphenylsulfonyl group,  
 4-chlorophenylcarbonyl group, 4-chlorophenylthiocarbonyl group,  
 4-fluorophenylcarbonyl group, 4-fluorophenylthiocarbonyl group,  
 isopropoxy carbonyl group, N-isopropylcarbamoyl group, N-isopropylthiocarbamoyl  
 group, butyloxy carbonyl group, N-butylcarbamoyl group, N-butylthiocarbamoyl  
 group, isobutyloxy carbonyl group, N-isobutylcarbamoyl group,  
 N-isobutylthiocarbamoyl group, t-butyloxy carbonyl group, N-t-butylcarbamoyl  
 group, N-t-butylthiocarbamoyl group, cyclopropyloxy carbonyl group,  
 N-cyclopropylcarbamoyl group, N-cyclopropylthiocarbamoyl group,  
 cyclopentyloxy carbonyl group, N-cyclopentylcarbamoyl group,

N-cyclopentylthiocarbamoyl group, cyclohexyloxycarbonyl group, N-cyclohexylcarbamoyl group, N-cyclohexylthiocarbamoyl group, cyclopentylmethyloxycarbonyl group, cyclohexylmethyloxycarbonyl group, phenyloxycarbonyl group, N-phenylcarbamoyl group, N-phenylthiocarbamoyl group, 4-methylphenyloxycarbonyl group, N-(4-methylphenyl)carbamoyl group, N-(4-methylphenyl)thiocarbamoyl group, 4-chlorophenyloxycarbonyl group, N-(4-chlorophenyl)carbamoyl group, N-(4-chlorophenyl)thiocarbamoyl group, 4-fluorophenyloxycarbonyl group, N-(4-fluorophenyl)carbamoyl group, N-(4-fluorophenyl)thiocarbamoyl group, (pyrrolidino-1-yl)carbonyl group, (piperidino-1-yl)carbonyl group, and (morpholino-4-yl)carbonyl group. Ry represents hydrogen atom, methyl group, ethyl group, or isobutyl group, or binds to Rz to form pyrrolidino group, piperidino group, piperazino group, morpholino group, pyrrol-1-yl group, imidazol-1-yl group, or pyrazol-1-yl group together with the nitrogen atom to which they binds.

AR represents naphthalen-2-yl group, naphthalen-1-yl group, benzofuran-5-yl group, benzofuran-4-yl group, benzofuran-2-yl group, benzo[b]thiophen-5-yl group, benzo[b]thiophen-4-yl group, benzo[b]thiophen-2-yl group, indol-5-yl group, indol-4-yl group, indol-6-yl group, benzothiazol-6-yl group, benzothiazol-7-yl group, benzothiazol-5-yl group, benzothiazol-4-yl group, dihydro-3H-benzothiazol-6-yl group, dihydro-3H-benzothiazol-7-yl group, dihydro-3H-benzothiazol-5-yl group, dihydro-3H-benzothiazol-4-yl group, quinolin-6-yl group, quinolin-3-yl group, quinolin-5-yl group, quinolin-7-yl group, dihydro-1H-quinolin-6-yl group, dihydro-1H-quinolin-5-yl group, benzo[d]isothiazol-5-yl group, benzo[d]isothiazol-4-yl group, benzo[d]isothiazol-6-yl group, benzo[d]isothiazol-7-yl group, 1H-indazol-5-yl group, 1H-indazol-4-yl group, 1H-indazol-6-yl group, benzo[c]isothiazol-5-yl group, benzo[c]isothiazol-4-yl group, benzo[c]isothiazol-6-yl group, benzo[c]isothiazol-7-yl group, 2H-indazol-5-yl group,

2H-indazol-4-yl group, 2H-indazol-6-yl group, imidazo[1,2-a]pyridin-6-yl group,  
 imidazo[1,2-a]pyridin-7-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1H-pyrrolo[2,3-b]pyridin-4-yl group, isoquinolin-6-yl group, isoquinolin-3-yl group,  
 isoquinolin-5-yl group, isoquinolin-7-yl group, dihydro-2H-isoquinolin-6-yl group,  
 dihydro-2H-isoquinolin-5-yl group, cinnolin-6-yl group, cinnolin-5-yl group,  
 quinazolin-6-yl group, quinazolin-7-yl group, quinazolin-5-yl group, quinoxalin-2-yl  
 group, quinoxalin-6-yl group, quinoxalin-5-yl group, 1H-benzimidazol-5-yl group,  
 1H-benzimidazol-4-yl group, benzoxazol-5-yl group, benzoxazol-6-yl group,  
 benzoxazol-4-yl group, benzoxazol-7-yl group, 1H-pyrrolo[3,2-b]pyridin-5-yl group,  
 1H-pyrrolo[3,2-b]pyridin-6-yl group, benzo[1,2,5]thiadiazol-5-yl group,  
 benzo[1,2,5]thiadiazol-4-yl group, 1H-benzotriazol-5-yl group, 1H-benzotriazol-4-yl  
 group, 1,3-dihydropyrrolo[2,3-b]pyridin-5-yl group,  
 1,3-dihydropyrrolo[2,3-b]pyridin-4-yl group, 1,3-dihydrobenzimidazol-5-yl group,  
 1,3-dihydrobenzimidazol-4-yl group, dihydro-3H-benzoxazol-6-yl group,  
 dihydro-3H-benzoxazol-7-yl group, dihydro-3H-benzoxazol-5-yl group,  
 dihydro-3H-benzoxazol-4-yl group, phthalazin-6-yl group, phthalazin-5-yl group,  
 [1,8]naphthalidin-3-yl group, [1,8]naphthalidin-4-yl group, [1,5]naphthalidin-3-yl  
 group, [1,5]naphthalidin-4-yl group, 1H-pyrrolo[3,2-c]pyridin-6-yl group,  
 1H-pyrrolo[3,2-c]pyridin-4-yl group, 1H-pyrrolo[2,3-c]pyridin-5-yl group,  
 1H-pyrrolo[2,3-c]pyridin-4-yl group, 1H-pyrazolo[4,3-b]pyridin-5-yl group,  
 1H-pyrazolo[4,3-b]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-6-yl group,  
 1H-pyrazolo[4,3-c]pyridin-4-yl group, 1H-pyrazolo[3,4-c]pyridin-5-yl group,  
 1H-pyrazolo[3,4-c]pyridin-4-yl group, 1H-pyrazolo[3,4-b]pyridin-5-yl group,  
 1H-pyrazolo[3,4-b]pyridin-4-yl group, [1,2,4]triazolo[4,3-a]pyridin-6-yl group,  
 [1,2,4]triazolo[4,3-a]pyridin-7-yl group, thieno[3,2-c]pyridin-2-yl group,  
 thieno[3,2-c]pyridin-3-yl group, thieno[3,2-c]pyridin-6-yl group,  
 thieno[3,2-b]pyridin-2-yl group, thieno[3,2-b]pyridin-3-yl group,

thieno[3,2-b]pyridin-5-yl group, thieno[3,2-b]pyridin-6-yl group, 1H-thieno[3,2-c]pyrazol-5-yl group, 1H-thieno[3,2-c]pyrazol-4-yl group, benzo[d]isoxazol-5-yl group, benzo[d]isoxazol-4-yl group, benzo[d]isoxazol-6-yl group, benzo[d]isoxazol-7-yl group, benzo[c]isoxazol-5-yl group, benzo[c]isoxazol-4-yl group, benzo[c]isoxazol-6-yl group, benzo[c]isoxazol-7-yl group, indolizin-7-yl group, indolizin-6-yl group, indolizine-8-yl group, 1,3-dihydroindol-5-yl group, 1,3-dihydroindol-4-yl group, 1,3-dihydroindol-6-yl group, 1H-pyrazolo[3,4-d]thiazol-5-yl group, 2H-isoindol-5-yl group, 2H-isoindol-4-yl group, [1,2,4]triazolo[1,5-a]pyrimidin-6-yl group, 1H-pyrazolo[3,4-b]pyrazin-5-yl group, 1H-imidazo[4,5-b]pyrazin-5-yl group, 7H-purin-2-yl group, 4H-chromen-6-yl group, or 4H-chromen-5-yl group (the aforementioned groups may be substituted with one of Xa or two or more of the same or different Xa). The substituent Xa represents a group as any one of oxo group, thioxo group, fluorine atom, chlorine atom, trifluoromethyl group, methyl group, ethyl group, propyl group, 2-hydroxyethyl group, carboxymethyl group, 2-carboxyethyl group, N,N-dimethylcarbamoylmethyl group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, carboxymethyloxy group, 2-carboxyethyloxy group, N,N-dimethylcarbamoylmethyloxy group, amino group, methylamino group, dimethylamino group, 2-hydroxyethylamino group, carbamoylamino group, acetylamino group, furan-2-carboxyamino group, 2-hydroxyacetylamino group, 2-aminoacetylamino group, methylsulfonylamino group, (N,N-dimethylsulfamoyl)amino group, methanesulfonyl group, sulfamoyl group, N-methylsulfamoyl group, N,N-dimethylsulfamoyl group, carboxyl group, acetyl group, carbamoyl group, and N,N-dimethylcarbamoyl group.

The group Y represents hydrogen atom, methyl group, or ethyl group.

In a preferred embodiment of the present invention, a compound or a salt thereof satisfying all of the following requirements is excluded from the compound represented by the formula (I) or a salt thereof.

Link represents  $-(CH_2)_n-$ , symbol n represents an integer of 1 to 3.

C<sup>3</sup> represents carbon atom to which AR bonds, C<sup>4</sup> represents carbon atom to which Rs bonds, C<sup>5</sup> represents a ring-constituting carbon atom which may be substituted with Zx, and C<sup>2</sup> and C<sup>6</sup> represent unsubstituted ring-constituting carbon atom.

Zx represents fluorine atom, chlorine atom, nitro group, amino group, methyl group, or a OR<sup>9</sup> group, and R<sup>9</sup> represents hydrogen atom or a lower alkyl group having 1 to 4 carbon atoms.

Rs represents -O-Rx. Rx represents a linear or branched saturated alkyl group having 3 to 8 carbon atoms, or represents Ra or Rb, Q in Rb represents a residue of a partially unsaturated or completely unsaturated monocyclic or condensed bicyclic carbon ring or heterocyclic ring (q), and binds to A<sup>2</sup> at an arbitrary position on the ring. The heterocyclic ring (q) contains one or two of the same or different ring-constituting heteroatoms selected from the group consisting of nitrogen atom, oxygen atom, and sulfur atom.

AR represents a residue of naphthalene, benzofuran, benzo[b]thiophene, indole, benzothiazole, dihydro-3H-benzothiazole, quinoline, dihydro-1H-quinoline, benzo[d]isothiazole, 1H-indazole, benzo[c]isothiazole, 2H-indazole, imidazo[1,2-a]pyridine, 1H-pyrrolo[2,3-b]pyridine, isoquinoline, or dihydro-2H-isoquinoline (the aforementioned residue may be substituted with one of Xa or two or more of the same or different Xa).

In another preferred embodiment of the present invention, the compound represented by the formula (I) or a salt thereof satisfies all of the following requirements.

Link represents  $-(CH_2)_n-$ , symbol n represents an integer of 1 to 3.

C<sup>3</sup> represents carbon atom to which AR bonds, C<sup>4</sup> represents carbon atom to which Rs bonds, C<sup>5</sup> may be replaced with V, and C<sup>2</sup> and C<sup>6</sup> represent unsubstituted

ring-constituting carbon atom.

V represents nitrogen atom, or carbon atom substituted with Zx, and Zx represents a group as any one of fluorine atom, chlorine atom, bromine atom, nitro group, methyl group, hydroxyl group, methoxy group, amino group, N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, and N,N-dimethylsulfamoylamino group.

Rs represents -O-Rx. Rx represents butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-cyclopentylethyl group, or 2-cyclohexylethyl group, or represents Rb or Rc. Q in Rb represents a group as any one of phenyl group, thienyl group, furyl group, pyridyl group, oxazolyl group, naphthyl group, tetrahydronaphthyl group, indanyl group, indolyl group, and dihydrobenzodioxyl group. A<sup>2</sup> represents a single bond, oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)- (provided that when A<sup>2</sup> represents oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)-, A<sup>1</sup> represents ethylene). R<sup>2</sup> and R<sup>3</sup> independently represent hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, dimethylamino group, acetylamino group, or methylsulfonylamino group (provided that when Q represents phenyl group, A<sup>1</sup> represents a single bond, or unsubstituted methylene, and A<sup>2</sup> represents a single bond, one of R<sup>2</sup> and R<sup>3</sup> represents a substituent other than hydrogen atom). Symbol p in Rc represents an integer of 2 or 3, and A<sup>4</sup> represents a single bond or methylene. A<sup>5</sup> represents -C(O)-, -C(S)-, or -S(O)<sub>2</sub>-. Rd represents hydrogen atom, or a group as any one of methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group, cyclopropylmethyl group, cyclopentyl group, cyclopentylmethyl group, cyclohexyl group, cyclohexylmethyl group, phenyl group,

4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, and pyridin-4-yl group. Re represents a group as any one of methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, phenylmethyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, furan-2-yl group, furan-3-yl group, thiophen-2-yl group, thiophen-3-yl group, methoxy group, ethoxy group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenyloxy group, 4-methylphenyloxy group, 4-chlorophenyloxy group, 4-fluorophenyloxy group, thiomethoxy group, amino group, N-methylamino group, N,N-dimethylamino group, N-ethylamino group, N,N-diethylamino group, N-propylamino group, N-isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group, N-(4-fluorophenyl)amino group, N-(pyridin-2-yl)amino group, N-(pyridin-3-yl)amino group, N-(pyridin-4-yl)amino group, N-(furan-2-yl)amino group, N-(furan-3-yl)amino group, N-(thiophen-2-yl)amino group, N-(thiophen-3-yl)amino group, pyrrolidino group, piperidino group, morpholino group, methyloxycarbonylamino group, and ethyloxycarbonylamino group.

AR represents any one of cinnolin-6-yl group, cinnolin-5-yl group, quinazolin-6-yl group, quinazolin-7-yl group, quinazolin-5-yl group, quinoxalin-2-yl



group, quinoxalin-6-yl group, quinoxalin-5-yl group, 1H-benzimidazol-5-yl group,  
 1H-benzimidazol-4-yl group, benzoxazol-5-yl group, benzoxazol-6-yl group,  
 benzoxazol-4-yl group, benzoxazol-7-yl group, 1H-pyrrolo[3,2-b]pyridin-5-yl group,  
 1H-pyrrolo[3,2-b]pyridin-6-yl group, benzo[1,2,5]thiadiazol-5-yl group,  
 benzo[1,2,5]thiadiazol-4-yl group, 1H-benzotriazol-5-yl group, 1H-benzotriazol-4-yl  
 group, 1,3-dihydropyrrolo[2,3-b]pyridin-5-yl group,  
 1,3-dihydropyrrolo[2,3-b]pyridin-4-yl group, 1,3-dihydrobenzimidazol-5-yl group,  
 1,3-dihydrobenzimidazol-4-yl group, dihydro-3H-benzoxazol-6-yl group,  
 dihydro-3H-benzoxazol-7-yl group, dihydro-3H-benzoxazol-5-yl group,  
 dihydro-3H-benzoxazol-4-yl group, phthalazin-6-yl group, phthalazin-5-yl group,  
 [1,8]naphthalidin-3-yl group, [1,8]naphthalidin-4-yl group, [1,5]naphthalidin-3-yl  
 group, [1,5]naphthalidin-4-yl group, 1H-pyrrolo[3,2-c]pyridin-6-yl group,  
 1H-pyrrolo[3,2-c]pyridin-4-yl group, 1H-pyrrolo[2,3-c]pyridin-5-yl group,  
 1H-pyrrolo[2,3-c]pyridin-4-yl group, 1H-pyrazolo[4,3-b]pyridin-5-yl group,  
 1H-pyrazolo[4,3-b]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-6-yl group,  
 1H-pyrazolo[4,3-c]pyridin-4-yl group, 1H-pyrazolo[3,4-c]pyridin-5-yl group,  
 1H-pyrazolo[3,4-c]pyridin-4-yl group, 1H-pyrazolo[3,4-b]pyridin-5-yl group,  
 1H-pyrazolo[3,4-b]pyridin-4-yl group, [1,2,4]triazolo[4,3-a]pyridin-6-yl group,  
 [1,2,4]triazolo[4,3-a]pyridin-7-yl group, thieno[3,2-c]pyridin-2-yl group,  
 thieno[3,2-c]pyridin-3-yl group, thieno[3,2-c]pyridin-6-yl group,  
 thieno[3,2-b]pyridin-2-yl group, thieno[3,2-b]pyridin-3-yl group,  
 thieno[3,2-b]pyridin-5-yl group, thieno[3,2-b]pyridin-6-yl group,  
 1H-thieno[3,2-c]pyrazol-5-yl group, 1H-thieno[3,2-c]pyrazol-4-yl group,  
 benzo[d]isoxazol-5-yl group, benzo[d]isoxazol-4-yl group, benzo[d]isoxazol-6-yl group,  
 benzo[d]isoxazol-7-yl group, benzo[c]isoxazol-5-yl group, benzo[c]isoxazol-4-yl group,  
 benzo[c]isoxazol-6-yl group, benzo[c]isoxazol-7-yl group, indolizin-7-yl group,  
 indolizin-6-yl group, indolizine-8-yl group, 1,3-dihydroindol-5-yl group,

1,3-dihydroindol-4-yl group, 1,3-dihydroindol-6-yl group, 1H-pyrazolo[3,4-d]thiazol-5-yl group, 2H-isoindol-5-yl group, 2H-isoindol-4-yl group, [1,2,4]triazolo[1,5-a]pyrimidin-6-yl group, 1H-pyrazolo[3,4-b]pyrazin-5-yl group, 1H-imidazo[4,5-b]pyrazin-5-yl group, 7H-purin-2-yl group, 4H-chromen-6-yl group, and 4H-chromen-5-yl group (these groups may be substituted with one of Xa or two or more of the same or different Xa). The substituent Xa represents a group as any one of oxo group, thioxo group, fluorine atom, chlorine atom, trifluoromethyl group, methyl group, ethyl group, propyl group, 2-hydroxyethyl group, carboxymethyl group, 2-carboxyethyl group, N,N-dimethylcarbamoylmethyl group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, carboxymethyloxy group, 2-carboxyethyloxy group, N,N-dimethylcarbamoylmethyloxy group, amino group, methylamino group, dimethylamino group, 2-hydroxyethylamino group, carbamoylamino group, acetylamino group, furan-2-carboxyamino group, 2-hydroxyacetylamino group, 2-aminoacetylamino group, methylsulfonylamino group, (N,N-dimethylsulfamoyl)amino group, methanesulfonyl group, sulfamoyl group, N-methylsulfamoyl group, N,N-dimethylsulfamoyl group, carboxyl group, acetyl group, carbamoyl group, and N,N-dimethylcarbamoyl group.

The group Y represents hydrogen atom, methyl group, or ethyl group.

In another preferred embodiment of the present invention, the compound represented by the formula (I) or a salt thereof satisfies all of the following requirements.

Link represents  $-(CH_2)_n-$ , symbol n represents an integer of 1 to 3.

C<sup>3</sup> represents carbon atom to which AR bonds, C<sup>4</sup> represents carbon atom to which Rs bonds, C<sup>5</sup> may be replaced with V, and C<sup>2</sup> and C<sup>6</sup> represent unsubstituted ring-constituting carbon atom.

V represents nitrogen atom, or carbon atom substituted with Zx, and Zx represents a group as any one of, chlorine atom, bromine atom, nitro group, methyl

group, hydroxyl group, methoxy group, amino group, N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, and N,N-dimethylsulfamoylamino group.

$R_s$  represents  $-S-R_x$ .  $R_x$  represents butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-cyclopentylethyl group, or 2-cyclohexylethyl group, or represents  $R_b$  or  $R_c$ .  $Q$  in  $R_b$  represents a group as any one of phenyl group, thienyl group, furyl group, pyridyl group, oxazolyl group, naphthyl group, tetrahydronaphthyl group, indanyl group, indolyl group, and dihydrobenzodioxyl group.  $A^2$  represents a single bond, oxygen atom, sulfur atom,  $-N(\text{methyl})-$ , or  $-N(\text{ethyl})-$  (provided that when  $A^2$  represents oxygen atom, sulfur atom,  $-N(\text{methyl})-$ , or  $-N(\text{ethyl})-$ ,  $A^1$  represents ethylene).  $R^2$  and  $R^3$  independently represent hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, dimethylamino group, acetylamino group, or methylsulfonylamino group (provided that when  $Q$  represents phenyl group,  $A^1$  represents a single bond, or unsubstituted methylene, and  $A^2$  represents a single bond, one of  $R^2$  and  $R^3$  represents a substituent other than hydrogen atom). Symbol  $p$  in  $R_c$  represents an integer of 2 or 3, and  $A^4$  represents a single bond or methylene.  $A^5$  represents  $-C(O)-$ ,  $-C(S)-$ , or  $-S(O)_2-$ .  $R_d$  represents hydrogen atom, or a group as any one of methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group, cyclopropylmethyl group, cyclopentyl group, cyclopentylmethyl group, cyclohexyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, and pyridin-4-yl group.  $R_e$  represents a group as any one of

methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, phenylmethyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, methoxy group, ethoxy group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenyloxy group, 4-methylphenyloxy group, 4-chlorophenyloxy group, 4-fluorophenyloxy group, thiomethoxy group, amino group, N-methylamino group, N,N-dimethylamino group, N-ethylamino group, N,N-diethylamino group, N-propylamino group, N-isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group, N-(4-fluorophenyl)amino group, N-(pyridin-2-yl)amino group, N-(pyridin-3-yl)amino group, N-(pyridin-4-yl)amino group, N-(furan-2-yl)amino group, N-(furan-3-yl)amino group, N-(thiophen-2-yl)amino group, N-(thiophen-3-yl)amino group, pyrrolidino group, piperidino group, morpholino group, methyloxycarbonylamino group, and ethyloxycarbonylamino group.

AR represents naphthalen-2-yl group, naphthalen-1-yl group, benzofuran-5-yl group, benzofuran-4-yl group, benzofuran-2-yl group, benzo[b]thiophen-5-yl group, benzo[b]thiophen-4-yl group, benzo[b]thiophen-2-yl group, indol-5-yl group, indol-4-yl group, indol-6-yl group, benzothiazol-6-yl group, benzothiazol-7-yl group, benzothiazol-5-yl group, benzothiazol-4-yl group, dihydro-3H-benzothiazol-6-yl group, dihydro-3H-benzothiazol-7-yl group,

dihydro-3H-benzothiazol-5-yl group, dihydro-3H-benzothiazol-4-yl group,  
 quinolin-6-yl group, quinolin-3-yl group, quinolin-5-yl group, quinolin-7-yl group,  
 dihydro-1H-quinolin-6-yl group, dihydro-1H-quinolin-5-yl group,  
 benzo[d]isothiazol-5-yl group, benzo[d]isothiazol-4-yl group, benzo[d]isothiazol-6-yl  
 group, benzo[d]isothiazol-7-yl group, 1H-indazol-5-yl group, 1H-indazol-4-yl group,  
 1H-indazol-6-yl group, benzo[c]isothiazol-5-yl group, benzo[c]isothiazol-4-yl group,  
 benzo[c]isothiazol-6-yl group, benzo[c]isothiazol-7-yl group, 2H-indazol-5-yl group,  
 2H-indazol-4-yl group, 2H-indazol-6-yl group, imidazo[1,2-a]pyridin-6-yl group,  
 imidazo[1,2-a]pyridin-7-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1H-pyrrolo[2,3-b]pyridin-4-yl group, isoquinolin-6-yl group, isoquinolin-3-yl group,  
 isoquinolin-5-yl group, isoquinolin-7-yl group, dihydro-2H-isoquinolin-6-yl group,  
 dihydro-2H-isoquinolin-5-yl group, cinnolin-6-yl group, cinnolin-5-yl group,  
 quinazolin-6-yl group, quinazolin-7-yl group, quinazolin-5-yl group, quinoxalin-2-yl  
 group, quinoxalin-6-yl group, quinoxalin-5-yl group, 1H-benzimidazol-5-yl group,  
 1H-benzimidazol-4-yl group, benzoxazol-5-yl group, benzoxazol-6-yl group,  
 benzoxazol-4-yl group, benzoxazol-7-yl group, 1H-pyrrolo[3,2-b]pyridin-5-yl group,  
 1H-pyrrolo[3,2-b]pyridin-6-yl group, benzo[1,2,5]thiadiazol-5-yl group,  
 benzo[1,2,5]thiadiazol-4-yl group, 1H-benzotriazol-5-yl group, 1H-benzotriazol-4-yl  
 group, 1,3-dihydropyrrolo[2,3-b]pyridin-5-yl group,  
 1,3-dihydropyrrolo[2,3-b]pyridin-4-yl group, 1,3-dihydrobenzimidazol-5-yl group,  
 1,3-dihydrobenzimidazol-4-yl group, dihydro-3H-benzoxazol-6-yl group,  
 dihydro-3H-benzoxazol-7-yl group, dihydro-3H-benzoxazol-5-yl group,  
 dihydro-3H-benzoxazol-4-yl group, phthalazin-6-yl group, phthalazin-5-yl group,  
 [1,8]naphthalidin-3-yl group, [1,8]naphthalidin-4-yl group, [1,5]naphthalidin-3-yl  
 group, [1,5]naphthalidin-4-yl group, 1H-pyrrolo[3,2-c]pyridin-6-yl group,  
 1H-pyrrolo[3,2-c]pyridin-4-yl group, 1H-pyrrolo[2,3-c]pyridin-5-yl group,  
 1H-pyrrolo[2,3-c]pyridin-4-yl group, 1H-pyrazolo[4,3-b]pyridin-5-yl group,

1H-pyrazolo[4,3-b]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-6-yl group,  
 1H-pyrazolo[4,3-c]pyridin-4-yl group, 1H-pyrazolo[3,4-c]pyridin-5-yl group,  
 1H-pyrazolo[3,4-c]pyridin-4-yl group, 1H-pyrazolo[3,4-b]pyridin-5-yl group,  
 1H-pyrazolo[3,4-b]pyridin-4-yl group, [1,2,4]triazolo[4,3-a]pyridin-6-yl group,  
 [1,2,4]triazolo[4,3-a]pyridin-7-yl group, thieno[3,2-c]pyridin-2-yl group,  
 thieno[3,2-c]pyridin-3-yl group, thieno[3,2-c]pyridin-6-yl group,  
 thieno[3,2-b]pyridin-2-yl group, thieno[3,2-b]pyridin-3-yl group,  
 thieno[3,2-b]pyridin-5-yl group, thieno[3,2-b]pyridin-6-yl group,  
 1H-thieno[3,2-c]pyrazol-5-yl group, 1H-thieno[3,2-c]pyrazol-4-yl group,  
 benzo[d]isoxazol-5-yl group, benzo[d]isoxazol-4-yl group, benzo[d]isoxazol-6-yl group,  
 benzo[d]isoxazol-7-yl group, benzo[c]isoxazol-5-yl group, benzo[c]isoxazol-4-yl group,  
 benzo[c]isoxazol-6-yl group, benzo[c]isoxazol-7-yl group, indolizin-7-yl group,  
 indolizin-6-yl group, indolizine-8-yl group, 1,3-dihydroindol-5-yl group,  
 1,3-dihydroindol-4-yl group, 1,3-dihydroindol-6-yl group,  
 1H-pyrazolo[3,4-d]thiazol-5-yl group, 2H-isindol-5-yl group, 2H-isindol-4-yl group,  
 [1,2,4]triazolo[1,5-a]pyrimidin-6-yl group, 1H-pyrazolo[3,4-b]pyrazin-5-yl group,  
 1H-imidazo[4,5-b]pyrazin-5-yl group, 7H-purin-2-yl group, 4H-chromen-6-yl group,  
 or 4H-chromen-5-yl group (these groups may be substituted with one of Xa or two or  
 more of the same or different Xa). The substituent Xa represents a group as any  
 one of oxo group, thioxo group, fluorine atom, chlorine atom, trifluoromethyl group,  
 methyl group, ethyl group, propyl group, 2-hydroxyethyl group, carboxymethyl  
 group, 2-carboxyethyl group, N,N-dimethylcarbamoylmethyl group, hydroxyl group,  
 methoxy group, 2-hydroxyethyloxy group, carboxymethyloxy group,  
 2-carboxyethyloxy group, N,N-dimethylcarbamoylmethyloxy group, amino group,  
 methylamino group, dimethylamino group, 2-hydroxyethylamino group,  
 carbamoylamino group, acetylamino group, furan-2-carboxyamino group,  
 2-hydroxyacetylamino group, 2-aminoacetylamino group, methylsulfonylamino

group, (N,N-dimethylsulfamoyl)amino group, methanesulfonyl group, sulfamoyl group, N-methylsulfamoyl group, N,N-dimethylsulfamoyl group, carboxyl group, acetyl group, carbamoyl group, and N,N-dimethylcarbamoyl group.

The group Y represents hydrogen atom, methyl group, or ethyl group.

In another preferred embodiment of the present invention, the compound represented by the formula (I) or a salt thereof satisfies all of the following requirements.

Link represents  $-(CH_2)_n-$ , symbol n represents an integer of 1 to 3.

C<sup>3</sup> represents carbon atom to which AR bonds, C<sup>4</sup> represents carbon atom to which Rs bonds, and C<sup>2</sup>, C<sup>5</sup> and C<sup>6</sup> represent unsubstituted ring-constituting carbon atom.

Rs represents  $-N(R_y)(R_z)$ . R<sub>z</sub> represents a group as any one of butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group,

2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, 2-(N-ethyl-N-phenylamino)ethyl group, isobutyryl group, isopropylthiocarbonyl group, isopropylsulfonyl group, valeryl group, butylthiocarbonyl group, isovaleryl group, isobutylthiocarbonyl group, pivaloyl group, t-butylthiocarbonyl group, cyclopropylcarbonyl group, cyclopropylthiocarbonyl group, cyclopentylcarbonyl group, cyclopentylthiocarbonyl group, cyclohexylcarbonyl group, cyclohexylthiocarbonyl group, cyclopentylmethylcarbonyl group, cyclopentylmethylthiocarbonyl group,



cyclohexylmethylcarbonyl group, cyclohexylmethylthiocarbonyl group, benzoyl group, thiobenzoyl group, phenylsulfonyl group, 4-methylphenylcarbonyl group, 4-methylphenylthiocarbonyl group, 4-methylphenylsulfonyl group, 4-chlorophenylcarbonyl group, 4-chlorophenylthiocarbonyl group, 4-fluorophenylcarbonyl group, 4-fluorophenylthiocarbonyl group, isopropoxy carbonyl group, N-isopropyl carbamoyl group, N-isopropylthiocarbamoyl group, butyloxy carbonyl group, N-butyl carbamoyl group, N-butylthiocarbamoyl group, isobutyloxy carbonyl group, N-isobutyl carbamoyl group, N-isobutylthiocarbamoyl group, t-butyloxy carbonyl group, N-t-butyl carbamoyl group, N-t-butylthiocarbamoyl group, cyclopropyloxy carbonyl group, N-cyclopropyl carbamoyl group, N-cyclopropylthiocarbamoyl group, cyclopentyloxy carbonyl group, N-cyclopentyl carbamoyl group, N-cyclopentylthiocarbamoyl group, cyclohexyloxy carbonyl group, N-cyclohexyl carbamoyl group, N-cyclohexylthiocarbamoyl group, cyclopentylmethyloxy carbonyl group, cyclohexylmethyloxy carbonyl group, phenyloxy carbonyl group, N-phenyl carbamoyl group, N-phenylthiocarbamoyl group, 4-methylphenyloxy carbonyl group, N-(4-methylphenyl) carbamoyl group, N-(4-methylphenyl)thiocarbamoyl group, 4-chlorophenyloxy carbonyl group, N-(4-chlorophenyl) carbamoyl group, N-(4-chlorophenyl)thiocarbamoyl group, 4-fluorophenyloxy carbonyl group, N-(4-fluorophenyl) carbamoyl group, N-(4-fluorophenyl)thiocarbamoyl group, (pyrrolidino-1-yl) carbonyl group, (piperidino-1-yl) carbonyl group, and (morpholino-4-yl) carbonyl group. Ry represents hydrogen atom, methyl group, ethyl group, or isobutyl group, or binds to Rz to form pyrrolidino group, piperidino group, piperazino group, morpholino group, pyrrol-1-yl group, imidazol-1-yl group, or pyrazol-1-yl group together with the nitrogen atom to which they bind.

AR represents naphthalen-2-yl group, naphthalen-1-yl group,

benzofuran-5-yl group, benzofuran-4-yl group, benzofuran-2-yl group,  
 benzo[b]thiophen-5-yl group, benzo[b]thiophen-4-yl group, benzo[b]thiophen-2-yl  
 group, indol-5-yl group, indol-4-yl group, indol-6-yl group, benzothiazol-6-yl group,  
 benzothiazol-7-yl group, benzothiazol-5-yl group, benzothiazol-4-yl group,  
 dihydro-3H-benzothiazol-6-yl group, dihydro-3H-benzothiazol-7-yl group,  
 dihydro-3H-benzothiazol-5-yl group, dihydro-3H-benzothiazol-4-yl group,  
 quinolin-6-yl group, quinolin-3-yl group, quinolin-5-yl group, quinolin-7-yl group,  
 dihydro-1H-quinolin-6-yl group, dihydro-1H-quinolin-5-yl group,  
 benzo[d]isothiazol-5-yl group, benzo[d]isothiazol-4-yl group, benzo[d]isothiazol-6-yl  
 group, benzo[d]isothiazol-7-yl group, 1H-indazol-5-yl group, 1H-indazol-4-yl group,  
 1H-indazol-6-yl group, benzo[c]isothiazol-5-yl group, benzo[c]isothiazol-4-yl group,  
 benzo[c]isothiazol-6-yl group, benzo[c]isothiazol-7-yl group, 2H-indazol-5-yl group,  
 2H-indazol-4-yl group, 2H-indazol-6-yl group, imidazo[1,2-a]pyridin-6-yl group,  
 imidazo[1,2-a]pyridin-7-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1H-pyrrolo[2,3-b]pyridin-4-yl group, isoquinolin-6-yl group, isoquinolin-3-yl group,  
 isoquinolin-5-yl group, isoquinolin-7-yl group, dihydro-2H-isoquinolin-6-yl group,  
 dihydro-2H-isoquinolin-5-yl group, cinnolin-6-yl group, cinnolin-5-yl group,  
 quinazolin-6-yl group, quinazolin-7-yl group, quinazolin-5-yl group, quinoxalin-2-yl  
 group, quinoxalin-6-yl group, quinoxalin-5-yl group, 1H-benzimidazol-5-yl group,  
 1H-benzimidazol-4-yl group, benzoxazol-5-yl group, benzoxazol-6-yl group,  
 benzoxazol-4-yl group, benzoxazol-7-yl group, 1H-pyrrolo[3,2-b]pyridin-5-yl group,  
 1H-pyrrolo[3,2-b]pyridin-6-yl group, benzo[1,2,5]thiadiazol-5-yl group,  
 benzo[1,2,5]thiadiazol-4-yl group, 1H-benzotriazol-5-yl group, 1H-benzotriazol-4-yl  
 group, 1,3-dihydropyrrolo[2,3-b]pyridin-5-yl group,  
 1,3-dihydropyrrolo[2,3-b]pyridin-4-yl group, 1,3-dihydrobenzimidazol-5-yl group,  
 1,3-dihydrobenzimidazol-4-yl group, dihydro-3H-benzoxazol-6-yl group,  
 dihydro-3H-benzoxazol-7-yl group, dihydro-3H-benzoxazol-5-yl group,

dihydro-3H-benzoxazol-4-yl group, phthalazin-6-yl group, phthalazin-5-yl group,  
 [1,8]naphthalidin-3-yl group, [1,8]naphthalidin-4-yl group, [1,5]naphthalidin-3-yl  
 group, [1,5]naphthalidin-4-yl group, 1H-pyrrolo[3,2-c]pyridin-6-yl group,  
 1H-pyrrolo[3,2-c]pyridin-4-yl group, 1H-pyrrolo[2,3-c]pyridin-5-yl group,  
 1H-pyrrolo[2,3-c]pyridin-4-yl group, 1H-pyrazolo[4,3-b]pyridin-5-yl group,  
 1H-pyrazolo[4,3-b]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-6-yl group,  
 1H-pyrazolo[4,3-c]pyridin-4-yl group, 1H-pyrazolo[3,4-c]pyridin-5-yl group,  
 1H-pyrazolo[3,4-c]pyridin-4-yl group, 1H-pyrazolo[3,4-b]pyridin-5-yl group,  
 1H-pyrazolo[3,4-b]pyridin-4-yl group, [1,2,4]triazolo[4,3-a]pyridin-6-yl group,  
 [1,2,4]triazolo[4,3-a]pyridin-7-yl group, thieno[3,2-c]pyridin-2-yl group,  
 thieno[3,2-c]pyridin-3-yl group, thieno[3,2-c]pyridin-6-yl group,  
 thieno[3,2-b]pyridin-2-yl group, thieno[3,2-b]pyridin-3-yl group,  
 thieno[3,2-b]pyridin-5-yl group, thieno[3,2-b]pyridin-6-yl group,  
 1H-thieno[3,2-c]pyrazol-5-yl group, 1H-thieno[3,2-c]pyrazol-4-yl group,  
 benzo[d]isoxazol-5-yl group, benzo[d]isoxazol-4-yl group, benzo[d]isoxazol-6-yl group,  
 benzo[d]isoxazol-7-yl group, benzo[c]isoxazol-5-yl group, benzo[c]isoxazol-4-yl group,  
 benzo[c]isoxazol-6-yl group, benzo[c]isoxazol-7-yl group, indolizin-7-yl group,  
 indolizin-6-yl group, indolizine-8-yl group, 1,3-dihydroindol-5-yl group,  
 1,3-dihydroindol-4-yl group, 1,3-dihydroindol-6-yl group,  
 1H-pyrazolo[3,4-d]thiazol-5-yl group, 2H-isoindol-5-yl group, 2H-isoindol-4-yl group,  
 [1,2,4]triazolo[1,5-a]pyrimidin-6-yl group, 1H-pyrazolo[3,4-b]pyrazin-5-yl group,  
 1H-imidazo[4,5-b]pyrazin-5-yl group, 7H-purin-2-yl group, 4H-chromen-6-yl group,  
 or 4H-chromen-5-yl group (these groups may be substituted with one of Xa or two or  
 more of the same or different Xa). The substituent Xa represents a group as any  
 one of oxo group, thioxo group, fluorine atom, chlorine atom, trifluoromethyl group,  
 methyl group, ethyl group, propyl group, 2-hydroxyethyl group, carboxymethyl  
 group, 2-carboxyethyl group, N,N-dimethylcarbamoylmethyl group, hydroxyl group,

methoxy group, 2-hydroxyethyloxy group, carboxymethyloxy group, 2-carboxyethyloxy group, N,N-dimethylcarbamoylethyloxy group, amino group, methylamino group, dimethylamino group, 2-hydroxyethylamino group, carbamoylethylamino group, acetylaminomethyl group, furan-2-carboxyaminomethyl group, 2-hydroxyacetylaminomethyl group, 2-aminoacetylaminomethyl group, methylsulfonylamino group, (N,N-dimethylsulfonylamino)amino group, methanesulfonyl group, sulfonyl group, N-methylsulfonyl group, N,N-dimethylsulfonyl group, carboxyl group, acetyl group, carbamoyl group, and N,N-dimethylcarbamoylethyl group.

The group Y represents hydrogen atom, methyl group, or ethyl group.

In another preferred embodiment of the present invention, the compound represented by the formula (I) or a salt thereof satisfies all of the following requirements.

Link represents  $-(CH_2)_n-$ , symbol n represents an integer of 1 to 3.

C<sup>3</sup> represents carbon atom to which AR bonds, C<sup>4</sup> represents carbon atom to which Rs bonds, C<sup>5</sup> may be replaced with V, and C<sup>2</sup> and C<sup>6</sup> represent unsubstituted ring-constituting carbon atom.

V represents nitrogen atom, or carbon atom substituted with Zx, and Zx represents a group as any one of chlorine atom, bromine atom, nitro group, methyl group, hydroxyl group, methoxy group, amino group, N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylaminomethyl group, carbamoylethylamino group, mesylamino group, and N,N-dimethylsulfonylamino group.

Rs represents  $-D-R_c$ , and D represents oxygen atom or sulfur atom. Symbol p in R<sub>c</sub> represents an integer of 2 or 3, and A<sup>4</sup> represents a single bond or methylene. A<sup>5</sup> represents  $-C(O)-$ ,  $-C(S)-$ , or  $-S(O)_2-$ . R<sub>d</sub> represents hydrogen atom, or a group as any one of methyl group, ethyl group, propyl group, isopropyl group, butyl group,

isobutyl group, cyclopropyl group, cyclopropylmethyl group, cyclopentyl group, cyclopentylmethyl group, cyclohexyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, and pyridin-4-yl group. Re represents a group as any one of methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, phenylmethyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, furan-2-yl group, furan-3-yl group, thiophen-2-yl group, thiophen-3-yl group, methoxy group, ethoxy group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenoxy group, 4-methylphenoxy group, 4-chlorophenoxy group, 4-fluorophenoxy group, thiomethoxy group, amino group, N-methylamino group, N,N-dimethylamino group, N-ethylamino group, N,N-diethylamino group, N-propylamino group, N-isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group, N-(4-fluorophenyl)amino group, N-(pyridin-2-yl)amino group, N-(pyridin-3-yl)amino group, N-(pyridin-4-yl)amino group, N-(furan-2-yl)amino group, N-(furan-3-yl)amino group, N-(thiophen-2-yl)amino group, N-(thiophen-3-yl)amino group, pyrrolidino group, piperidino group, morpholino group, methyloxycarbonylamino group, and ethyloxycarbonylamino group.

AR represents naphthalen-2-yl group, naphthalen-1-yl group, benzofuran-5-yl group, benzofuran-4-yl group, benzofuran-2-yl group, benzo[b]thiophen-5-yl group, benzo[b]thiophen-4-yl group, benzo[b]thiophen-2-yl group, indol-5-yl group, indol-4-yl group, indol-6-yl group, benzothiazol-6-yl group, benzothiazol-7-yl group, benzothiazol-5-yl group, benzothiazol-4-yl group, dihydro-3H-benzothiazol-6-yl group, dihydro-3H-benzothiazol-7-yl group, dihydro-3H-benzothiazol-5-yl group, dihydro-3H-benzothiazol-4-yl group, quinolin-6-yl group, quinolin-3-yl group, quinolin-5-yl group, quinolin-7-yl group, dihydro-1H-quinolin-6-yl group, dihydro-1H-quinolin-5-yl group, benzo[d]isothiazol-5-yl group, benzo[d]isothiazol-4-yl group, benzo[d]isothiazol-6-yl group, benzo[d]isothiazol-7-yl group, 1H-indazol-5-yl group, 1H-indazol-4-yl group, 1H-indazol-6-yl group, benzo[c]isothiazol-5-yl group, benzo[c]isothiazol-4-yl group, benzo[c]isothiazol-6-yl group, benzo[c]isothiazol-7-yl group, 2H-indazol-5-yl group, 2H-indazol-4-yl group, 2H-indazol-6-yl group, imidazo[1,2-a]pyridin-6-yl group, imidazo[1,2-a]pyridin-7-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1H-pyrrolo[2,3-b]pyridin-4-yl group, isoquinolin-6-yl group, isoquinolin-3-yl group, isoquinolin-5-yl group, isoquinolin-7-yl group, dihydro-2H-isoquinolin-6-yl group, dihydro-2H-isoquinolin-5-yl group, cinnolin-6-yl group, cinnolin-5-yl group, quinazolin-6-yl group, quinazolin-7-yl group, quinazolin-5-yl group, quinoxalin-2-yl group, quinoxalin-6-yl group, quinoxalin-5-yl group, 1H-benzimidazol-5-yl group, 1H-benzimidazol-4-yl group, benzoxazol-5-yl group, benzoxazol-6-yl group, benzoxazol-4-yl group, benzoxazol-7-yl group, 1H-pyrrolo[3,2-b]pyridin-5-yl group, 1H-pyrrolo[3,2-b]pyridin-6-yl group, benzo[1,2,5]thiadiazol-5-yl group, benzo[1,2,5]thiadiazol-4-yl group, 1H-benzotriazol-5-yl group, 1H-benzotriazol-4-yl group, 1,3-dihydropyrrolo[2,3-b]pyridin-5-yl group, 1,3-dihydropyrrolo[2,3-b]pyridin-4-yl group, 1,3-dihydrobenzimidazol-5-yl group, 1,3-dihydrobenzimidazol-4-yl group, dihydro-3H-benzoxazol-6-yl group,

dihydro-3H-benzoxazol-7-yl group, dihydro-3H-benzoxazol-5-yl group,  
 dihydro-3H-benzoxazol-4-yl group, phthalazin-6-yl group, phthalazin-5-yl group,  
 [1,8]naphthalidin-3-yl group, [1,8]naphthalidin-4-yl group, [1,5]naphthalidin-3-yl  
 group, [1,5]naphthalidin-4-yl group, 1H-pyrrolo[3,2-c]pyridin-6-yl group,  
 1H-pyrrolo[3,2-c]pyridin-4-yl group, 1H-pyrrolo[2,3-c]pyridin-5-yl group,  
 1H-pyrrolo[2,3-c]pyridin-4-yl group, 1H-pyrazolo[4,3-b]pyridin-5-yl group,  
 1H-pyrazolo[4,3-b]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-6-yl group,  
 1H-pyrazolo[4,3-c]pyridin-4-yl group, 1H-pyrazolo[3,4-c]pyridin-5-yl group,  
 1H-pyrazolo[3,4-c]pyridin-4-yl group, 1H-pyrazolo[3,4-b]pyridin-5-yl group,  
 1H-pyrazolo[3,4-b]pyridin-4-yl group, [1,2,4]triazolo[4,3-a]pyridin-6-yl group,  
 [1,2,4]triazolo[4,3-a]pyridin-7-yl group, thieno[3,2-c]pyridin-2-yl group,  
 thieno[3,2-c]pyridin-3-yl group, thieno[3,2-c]pyridin-6-yl group,  
 thieno[3,2-b]pyridin-2-yl group, thieno[3,2-b]pyridin-3-yl group,  
 thieno[3,2-b]pyridin-5-yl group, thieno[3,2-b]pyridin-6-yl group,  
 1H-thieno[3,2-c]pyrazol-5-yl group, 1H-thieno[3,2-c]pyrazol-4-yl group,  
 benzo[d]isoxazol-5-yl group, benzo[d]isoxazol-4-yl group, benzo[d]isoxazol-6-yl group,  
 benzo[d]isoxazol-7-yl group, benzo[c]isoxazol-5-yl group, benzo[c]isoxazol-4-yl group,  
 benzo[c]isoxazol-6-yl group, benzo[c]isoxazol-7-yl group, indolizin-7-yl group,  
 indolizin-6-yl group, indolizine-8-yl group, 1,3-dihydroindol-5-yl group,  
 1,3-dihydroindol-4-yl group, 1,3-dihydroindol-6-yl group,  
 1H-pyrazolo[3,4-d]thiazol-5-yl group, 2H-isoindol-5-yl group, 2H-isoindol-4-yl group,  
 [1,2,4]triazolo[1,5-a]pyrimidin-6-yl group, 1H-pyrazolo[3,4-b]pyrazin-5-yl group,  
 1H-imidazo[4,5-b]pyrazin-5-yl group, 7H-purin-2-yl group, 4H-chromen-6-yl group,  
 or 4H-chromen-5-yl group (these groups may be substituted with one of Xa or two or  
 more of the same or different Xa). The substituent Xa represents a group as any  
 one of oxo group, thioxo group, fluorine atom, chlorine atom, trifluoromethyl group,  
 methyl group, ethyl group, propyl group, 2-hydroxyethyl group, carboxymethyl

group, 2-carboxyethyl group, N,N-dimethylcarbamoylmethyl group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, carboxymethyloxy group, 2-carboxyethyloxy group, N,N-dimethylcarbamoylmethyloxy group, amino group, methylamino group, dimethylamino group, 2-hydroxyethylamino group, carbamoylamino group, acetylamino group, furan-2-carboxyamino group, 2-hydroxyacetylamino group, 2-aminoacetylamino group, methylsulfonylamino group, (N,N-dimethylsulfamoyl)amino group, methanesulfonyl group, sulfamoyl group, N-methylsulfamoyl group, N,N-dimethylsulfamoyl group, carboxyl group, acetyl group, carbamoyl group, and N,N-dimethylcarbamoyl group.

The group Y represents hydrogen atom, methyl group, or ethyl group.

In another preferred embodiment of the present invention, the compound represented by the formula (I) or a salt thereof satisfies all of the following requirements.

Link is  $-(CH_2)_n-$ , n is an integer of 1 to 3, C<sup>3</sup> is carbon atom bound with AR, C<sup>4</sup> is carbon atom bound with Rs, C<sup>5</sup> may be replaced with V, C<sup>2</sup> and C<sup>6</sup> are unsubstituted ring-constituting carbon atoms,

V is nitrogen atom or V is carbon atom substituted with Zx, Zx is any one of fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, or N,N-dimethylamino group,

Rs is -D-Rx, D is a single bond, Rx is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-cyclopentylethyl group, or 2-cyclohexylethyl group, or Rx is Rb or Rc (provided that Q in Rb is phenyl group, thienyl group, furyl group, pyridyl group, oxazolyl group, naphthyl group, tetrahydronaphthyl group, indanyl group, indolyl group, or dihydrobenzodioxyl group), A<sup>2</sup> is a single bond, oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)- (provided that when A<sup>2</sup> represents oxygen atom, sulfur atom, -N(methyl)- or



$\cdot\text{N(ethyl)}\cdot$ ,  $\text{A}^1$  represents ethylene),  $\text{R}^2$  and  $\text{R}^3$  independently represent hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, dimethylamino group, acetylamino group, or methylsulfonylamino group,  $p$  in  $\text{Rc}$  is an integer of 2 or 3,  $\text{A}^4$  is a single bond or methylene,  $\text{A}^5$  is  $\cdot\text{C(O)}\cdot$ ,  $\cdot\text{C(S)}\cdot$ , or  $\cdot\text{S(O)}_2\cdot$ ,  $\text{Rd}$  is hydrogen atom, or methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group, cyclopropylmethyl group, cyclopentyl group, cyclopentylmethyl group, cyclohexyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, or pyridin-4-yl group,  $\text{Re}$  is methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, *t*-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, phenylmethyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, methoxy group, ethoxy group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, *t*-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenyloxy group, 4-methylphenyloxy group, 4-chlorophenyloxy group, 4-fluorophenyloxy group, thiomethoxy group, amino group, *N*-methylamino group, *N,N*-dimethylamino group, *N*-ethylamino group, *N,N*-diethylamino group, *N*-propylamino group, *N*-isopropylamino group, *N*-butylamino group, *N*-isobutylamino group, *N*-*t*-butylamino group, *N*-cyclopropylamino group, *N*-cyclopentylamino group, *N*-cyclohexylamino group, *N*-phenylamino group, *N*-(4-methylphenyl)amino group, *N*-(4-chlorophenyl)amino group, *N*-(4-fluorophenyl)amino group, *N*-(pyridin-2-yl)amino group, *N*-(pyridin-3-yl)amino group, *N*-(pyridin-4-yl)amino group, *N*-(furan-2-yl)amino group, *N*-(furan-3-yl)amino

group, N-(thiophen-2-yl)amino group, N-(thiophen-3-yl)amino group, pyrrolidino group, piperidino group, morpholino group, methyloxycarbonylamino group, or ethyloxycarbonylamino group,

AR is naphthalen-2-yl group, naphthalen-1-yl group, benzofuran-5-yl group, benzofuran-4-yl group, benzofuran-2-yl group, benzo[b]thiophen-5-yl group, benzo[b]thiophen-4-yl group, benzo[b]thiophen-2-yl group, indol-5-yl group, indol-4-yl group, indol-6-yl group, benzothiazol-6-yl group, benzothiazol-7-yl group, benzothiazol-5-yl group, benzothiazol-4-yl group, dihydro-3H-benzothiazol-6-yl group, dihydro-3H-benzothiazol-7-yl group, dihydro-3H-benzothiazol-5-yl group, dihydro-3H-benzothiazol-4-yl group, quinolin-6-yl group, quinolin-3-yl group, quinolin-5-yl group, quinolin-7-yl group, dihydro-1H-quinolin-6-yl group, dihydro-1H-quinolin-5-yl group, benzo[d]isothiazol-5-yl group, benzo[d]isothiazol-4-yl group, benzo[d]isothiazol-6-yl group, benzo[d]isothiazol-7-yl group, 1H-indazol-5-yl group, 1H-indazol-4-yl group, 1H-indazol-6-yl group, benzo[c]isothiazol-5-yl group, benzo[c]isothiazol-4-yl group, benzo[c]isothiazol-6-yl group, benzo[c]isothiazol-7-yl group, 2H-indazol-5-yl group, 2H-indazol-4-yl group, 2H-indazol-6-yl group, imidazo[1,2-a]pyridin-6-yl group, imidazo[1,2-a]pyridin-7-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1H-pyrrolo[2,3-b]pyridin-4-yl group, isoquinolin-6-yl group, isoquinolin-3-yl group, isoquinolin-5-yl group, isoquinolin-7-yl group, dihydro-2H-isoquinolin-6-yl group, dihydro-2H-isoquinolin-5-yl group, cinnolin-6-yl group, cinnolin-5-yl group, quinazolin-6-yl group, quinazolin-7-yl group, quinazolin-5-yl group, quinoxalin-2-yl group, quinoxalin-6-yl group, quinoxalin-5-yl group, 1H-benzimidazol-5-yl group, 1H-benzimidazol-4-yl group, benzoxazol-5-yl group, benzoxazol-6-yl group, benzoxazol-4-yl group, benzoxazol-7-yl group, 1H-pyrrolo[3,2-b]pyridin-5-yl group, 1H-pyrrolo[3,2-b]pyridin-6-yl group, benzo[1,2,5]thiadiazol-5-yl group, benzo[1,2,5]thiadiazol-4-yl group, 1H-benzotriazol-5-yl group, 1H-benzotriazol-4-yl

group, 1,3-dihydropyrrolo[2,3-b]pyridin-5-yl group,  
 1,3-dihydropyrrolo[2,3-b]pyridin-4-yl group, 1,3-dihydrobenzimidazol-5-yl group,  
 1,3-dihydrobenzimidazol-4-yl group, dihydro-3H-benzoxazol-6-yl group,  
 dihydro-3H-benzoxazol-7-yl group, dihydro-3H-benzoxazol-5-yl group,  
 dihydro-3H-benzoxazol-4-yl group, phthalazin-6-yl group, phthalazin-5-yl group,  
 [1,8]naphthalidin-3-yl group, [1,8]naphthalidin-4-yl group, [1,5]naphthalidin-3-yl  
 group, [1,5]naphthalidin-4-yl group, 1H-pyrrolo[3,2-c]pyridin-6-yl group,  
 1H-pyrrolo[3,2-c]pyridin-4-yl group, 1H-pyrrolo[2,3-c]pyridin-5-yl group,  
 1H-pyrrolo[2,3-c]pyridin-4-yl group, 1H-pyrazolo[4,3-b]pyridin-5-yl group,  
 1H-pyrazolo[4,3-b]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-6-yl group,  
 1H-pyrazolo[4,3-c]pyridin-4-yl group, 1H-pyrazolo[3,4-c]pyridin-5-yl group,  
 1H-pyrazolo[3,4-c]pyridin-4-yl group, 1H-pyrazolo[3,4-b]pyridin-5-yl group,  
 1H-pyrazolo[3,4-b]pyridin-4-yl group, [1,2,4]triazolo[4,3-a]pyridin-6-yl group,  
 [1,2,4]triazolo[4,3-a]pyridin-7-yl group, thieno[3,2-c]pyridin-2-yl group,  
 thieno[3,2-c]pyridin-3-yl group, thieno[3,2-c]pyridin-6-yl group,  
 thieno[3,2-b]pyridin-2-yl group, thieno[3,2-b]pyridin-3-yl group,  
 thieno[3,2-b]pyridin-5-yl group, thieno[3,2-b]pyridin-6-yl group,  
 1H-thieno[3,2-c]pyrazol-5-yl group, 1H-thieno[3,2-c]pyrazol-4-yl group,  
 benzo[d]isoxazol-5-yl group, benzo[d]isoxazol-4-yl group, benzo[d]isoxazol-6-yl group,  
 benzo[d]isoxazol-7-yl group, benzo[c]isoxazol-5-yl group, benzo[c]isoxazol-4-yl group,  
 benzo[c]isoxazol-6-yl group, benzo[c]isoxazol-7-yl group, indolizin-7-yl group,  
 indolizin-6-yl group, indolizine-8-yl group, 1,3-dihydroindol-5-yl group,  
 1,3-dihydroindol-4-yl group, 1,3-dihydroindol-6-yl group,  
 1H-pyrazolo[3,4-d]thiazol-5-yl group, 2H-isoindol-5-yl group, 2H-isoindol-4-yl group,  
 [1,2,4]triazolo[1,5-a]pyrimidin-6-yl group, 1H-pyrazolo[3,4-b]pyrazin-5-yl group,  
 1H-imidazo[4,5-b]pyrazin-5-yl group, 7H-purin-2-yl group, 4H-chromen-6-yl group,  
 or 4H-chromen-5-yl group (the aforementioned groups may be substituted with one

of Xa or two or more of the same or different Xa), Xa is oxo group, thioxo group, fluorine atom, chlorine atom, trifluoromethyl group, methyl group, ethyl group, propyl group, 2-hydroxyethyl group, carboxymethyl group, 2-carboxyethyl group, N,N-dimethylcarbamoylemethyl group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, carboxymethyloxy group, 2-carboxyethyloxy group, N,N-dimethylcarbamoylemethyloxy group, amino group, methylamino group, dimethylamino group, 2-hydroxyethylamino group, carbamoylamino group, acetylamino group, furan-2-carboxyamino group, 2-hydroxyacetylamino group, 2-aminoacetylamino group, methylsulfonylamino group, (N,N-dimethylsulfamoyl)amino group, methanesulfonyl group, sulfamoyl group, N-methylsulfamoyl group, N,N-dimethylsulfamoyl group, carboxyl group, acetyl group, carbamoyl group, or N,N-dimethylcarbamoylemethyl group, and

Y is hydrogen atom, methyl group, or ethyl group.

In a particularly preferred embodiment of the present invention, the compound represented by the formula (I) or a salt thereof satisfies all of the following requirements.

Link represents  $-(CH_2)_n-$ , symbol n represents an integer of 2.

C<sup>2</sup> represents carbon atom to which AR bonds, C<sup>3</sup> represents carbon atom to which Rs bonds, C<sup>4</sup> may be replaced with V, and C<sup>5</sup> and C<sup>6</sup> represent unsubstituted ring-constituting carbon atom.

V represents nitrogen atom, or carbon atom substituted with Zx, and Zx represents a group as any one of fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, and N,N-dimethylamino group.

Rs represents -O-Rx. Rx represents a group as any one of butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl

group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl  
 group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl  
 group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl  
 group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl  
 group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group,  
 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl  
 group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group,  
 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group,  
 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group,  
 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl  
 group, 3-methylphenylmethyl group, 4-methylphenylmethyl group,  
 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group,  
 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl  
 group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group,  
 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group,  
 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group,  
 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group,  
 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group,  
 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group,  
 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group,  
 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group,  
 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group,  
 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group,  
 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group,  
 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group,  
 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,  
 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,

2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group,  
 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group,  
 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group,  
 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group,  
 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group,  
 2-(N-phenyl-N-methylamino)ethyl group, and 2-(N-ethyl-N-phenylamino)ethyl  
 group.

AR represents a group as any one of naphthalen-2-yl group,  
 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group,  
 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group,  
 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl  
 group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group,  
 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group,  
 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group,  
 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group,  
 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group,  
 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group,  
 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group,  
 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group,  
 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group,  
 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group,  
 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group,  
 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group,  
 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group,  
 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group,

2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, and benzoxazol-5-yl group.

The group Y represents hydrogen atom, methyl group, or ethyl group.

In another particularly preferred embodiment of the present invention, the compound represented by the formula (I) or a salt thereof satisfies all of the following requirements.

Link represents  $-(CH_2)_n$ , symbol n represents an integer of 2.

C<sup>2</sup> represents carbon atom to which AR bonds, C<sup>4</sup> represents carbon atom to which Rs bonds, C<sup>5</sup> may be replaced with V, and C<sup>3</sup> and C<sup>6</sup> represent unsubstituted ring-constituting carbon atom.

V represents nitrogen atom, or carbon atom substituted with Zx, and Zx represents a group as any one of fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, and N,N-dimethylamino group.

Rs represents -O-Rx. Rx represents a group as any one of butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group,



2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group,  
 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group,  
 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,  
 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,  
 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group,  
 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group,  
 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group,  
 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group,  
 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group,  
 2-(N-phenyl-N-methylamino)ethyl group, and 2-(N-ethyl-N-phenylamino)ethyl group.

AR represents a group as any one of naphthalen-2-yl group,  
 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group,  
 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group,  
 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group,  
 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group,  
 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group,  
 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group,  
 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group,  
 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group,  
 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group,  
 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group,  
 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group,  
 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group,  
 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group,  
 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group,  
 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group,

2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group,  
 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group,  
 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group,  
 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group,  
 quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl  
 group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group,  
 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group,  
 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group,  
 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group,  
 imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl  
 group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group,  
 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, and benzoxazol-5-yl  
 group.

The group Y represents hydrogen atom, methyl group, or ethyl group.

In another particularly preferred embodiment of the present invention, the  
 compound represented by the formula (I) or a salt thereof satisfies all of the  
 following requirements.

Link represents  $-(CH_2)_n-$ , symbol n represents an integer of 2.

C<sup>3</sup> represents carbon atom to which AR bonds, C<sup>5</sup> represents carbon atom to which Rs bonds, and C<sup>2</sup>, C<sup>4</sup> and C<sup>6</sup> represent unsubstituted ring-constituting carbon atom.

Rs represents -O-Rx. Rx represents a group as any one of butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group,

2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, and 2-(N-ethyl-N-phenylamino)ethyl group.

AR represents a group as any one of naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group,

1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group, 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, and benzoxazol-5-yl group.

The group Y represents hydrogen atom, methyl group, or ethyl group.

In another particularly preferred embodiment of the present invention, the compound represented by the formula (I) or a salt thereof satisfies all of the following requirements.

Link represents  $-(CH_2)_n-$ , symbol  $n$  represents an integer of 2.

$C^3$  represents carbon atom to which AR bonds,  $C^4$  represents carbon atom to which Rs bonds,  $C^5$  represents nitrogen atom, and  $C^2$  and  $C^6$  represent unsubstituted ring-constituting carbon atom.

Rs represents  $-O-Rx$ . Rx represents a group as any one of butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group,

3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group,  
 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group,  
 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group,  
 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group,  
 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group,  
 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group,  
 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,  
 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,  
 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group,  
 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group,  
 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group,  
 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group,  
 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group,  
 2-(N-phenyl-N-methylamino)ethyl group, and 2-(N-ethyl-N-phenylamino)ethyl group.

AR represents a group as any one of naphthalen-2-yl group,  
 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group,  
 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group,  
 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group,  
 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group,  
 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group,  
 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group,  
 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group,  
 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group,  
 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group,  
 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group,  
 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group,

1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group,  
 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group,  
 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group,  
 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group,  
 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group,  
 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group,  
 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group,  
 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group,  
 quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl  
 group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group,  
 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group,  
 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group,  
 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group,  
 imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl  
 group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group,  
 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, and benzoxazol-5-yl  
 group.

The group Y represents hydrogen atom, methyl group, or ethyl group.

In another preferred embodiment of the present invention, the compound  
 represented by the formula (I) or a salt thereof satisfies all of the following



requirements.

Link represents  $-(CH_2)_n-$ , symbol  $n$  represents an integer of 2.

$C^3$  represents carbon atom to which AR bonds,  $C^4$  represents carbon atom to which Rs bonds,  $C^6$  represents carbon atom substituted with Zx, and  $C^2$  and  $C^5$  represent unsubstituted ring-constituting carbon atom.

Zx represents fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, or N,N-dimethylamino group.

Rs represents -O-Rx. Rx represents a group as any one of butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group,

3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group,  
 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group,  
 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group,  
 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group,  
 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group,  
 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group,  
 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group,  
 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group,  
 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group,  
 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,  
 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,  
 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group,  
 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group,  
 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group,  
 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group,  
 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group,  
 2-(N-phenyl-N-methylamino)ethyl group, and 2-(N-ethyl-N-phenylamino)ethyl  
 group.

AR represents a group as any one of naphthalen-2-yl group,  
 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group,  
 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group,  
 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl  
 group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group,  
 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group,  
 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group,  
 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group,  
 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group,

2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group,  
 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group,  
 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group,  
 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group,  
 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group,  
 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group,  
 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group,  
 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group,  
 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group,  
 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group,  
 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group,  
 quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl  
 group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group,  
 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group,  
 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group,  
 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group,  
 imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl  
 group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group,  
 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, and benzoxazol-5-yl  
 group.

The group Y represents hydrogen atom, methyl group, or ethyl group.

In another particularly preferred embodiment of the present invention, the compound represented by the formula (I) or a salt thereof satisfies all of the following requirements.

Link represents  $-(CH_2)_n-$ , symbol n represents an integer of 2.

C<sup>3</sup> represents carbon atom to which AR bonds, C<sup>4</sup> represents carbon atom to which Rs bonds, and C<sup>2</sup>, C<sup>5</sup> and C<sup>6</sup> represent unsubstituted ring-constituting carbon atom.

Rs represents  $-N(R_y)(R_z)$ . R<sub>z</sub> represents a group as any one of butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group,

2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group,  
 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group,  
 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group,  
 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group,  
 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group,  
 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group,  
 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group,  
 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group,  
 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group,  
 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group,  
 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,  
 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,  
 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group,  
 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group,  
 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group,  
 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group,  
 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group,  
 2-(N-phenyl-N-methylamino)ethyl group, 2-(N-ethyl-N-phenylamino)ethyl group,  
 isobutyryl group, isopropylthiocarbonyl group, isopropylsulfonyl group, valeryl  
 group, butylthiocarbonyl group, isovaleryl group, isobutylthiocarbonyl group,  
 pivaloyl group, t-butylthiocarbonyl group, cyclopropylcarbonyl group,  
 cyclopropylthiocarbonyl group, cyclopentylcarbonyl group, cyclopentylthiocarbonyl  
 group, cyclohexylcarbonyl group, cyclohexylthiocarbonyl group,  
 cyclopentylmethylcarbonyl group, cyclopentylmethylthiocarbonyl group,  
 cyclohexylmethylcarbonyl group, cyclohexylmethylthiocarbonyl group, benzoyl  
 group, thiobenzoyl group, phenylsulfonyl group, 4-methylphenylcarbonyl group,  
 4-methylphenylthiocarbonyl group, 4-methylphenylsulfonyl group,

4-chlorophenylcarbonyl group, 4-chlorophenylthiocarbonyl group,  
 4-fluorophenylcarbonyl group, 4-fluorophenylthiocarbonyl group,  
 isopropylloxycarbonyl group, N-isopropylcarbamoyl group, N-isopropylthiocarbamoyl  
 group, butylloxycarbonyl group, N-butylcarbamoyl group, N-butylthiocarbamoyl  
 group, isobutylloxycarbonyl group, N-isobutylcarbamoyl group,  
 N-isobutylthiocarbamoyl group, t-butylloxycarbonyl group, N-t-butylcarbamoyl  
 group, N-t-butylthiocarbamoyl group, cyclopropylloxycarbonyl group,  
 N-cyclopropylcarbamoyl group, N-cyclopropylthiocarbamoyl group,  
 cyclopentylloxycarbonyl group, N-cyclopentylcarbamoyl group,  
 N-cyclopentylthiocarbamoyl group, cyclohexylloxycarbonyl group,  
 N-cyclohexylcarbamoyl group, N-cyclohexylthiocarbamoyl group,  
 cyclopentylmethyloxycarbonyl group, cyclohexylmethyloxycarbonyl group,  
 phenylloxycarbonyl group, N-phenylcarbamoyl group, N-phenylthiocarbamoyl group,  
 4-methylphenylloxycarbonyl group, N-(4-methylphenyl)carbamoyl group,  
 N-(4-methylphenyl)thiocarbamoyl group, 4-chlorophenylloxycarbonyl group,  
 N-(4-chlorophenyl)carbamoyl group, N-(4-chlorophenyl)thiocarbamoyl group,  
 4-fluorophenylloxycarbonyl group, N-(4-fluorophenyl)carbamoyl group,  
 N-(4-fluorophenyl)thiocarbamoyl group, (pyrrolidino-1-yl)carbonyl group,  
 (piperidino-1-yl)carbonyl group, and (morpholino-4-yl)carbonyl group. R<sub>y</sub>  
 represents hydrogen atom, methyl group, ethyl group, or isobutyl group, or binds to  
 R<sub>z</sub> to form pyrrolidino group, piperidino group, or morpholino group together with  
 nitrogen atom to which they bonds.

AR represents a group as any one of naphthalen-2-yl group,  
 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group,  
 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group,  
 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl  
 group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group,

2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group,  
 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group,  
 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group,  
 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group,  
 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group,  
 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group,  
 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group,  
 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group,  
 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group,  
 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group,  
 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group,  
 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group,  
 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group,  
 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group,  
 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group,  
 quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl  
 group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group,  
 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group,  
 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group,  
 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group,  
 imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl

group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group,  
1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, and benzoxazol-5-yl  
group.

The group Y represents hydrogen atom, methyl group, or ethyl group.

In another particularly preferred embodiment of the present invention, the  
compound represented by the formula (I) or a salt thereof satisfies all of the  
following requirements.

Link represents  $-(CH_2)_n-$ , symbol n represents an integer of 2.

C<sup>3</sup> represents carbon atom to which AR bonds, C<sup>4</sup> represents carbon atom to  
which Rs bonds, and C<sup>2</sup>, C<sup>5</sup> and C<sup>6</sup> represent unsubstituted ring-constituting carbon  
atom.

Rs represents -N(Ry)(Rz). -N(Ry)(Rz) is any one of N,N-dimethylamino  
group, N-ethyl-N-methylamino group, N,N-diethylamino group,  
N-methyl-N-propylamino group, N-ethyl-N-propylamino group,  
N-isopropyl-N-methylamino group, N-ethyl-N-isopropylamino group, N-butylamino  
group, N-butyl-N-methylamino group, N-butyl-N-ethylamino group,  
N-isobutylamino group, N-isobutyl-N-methylamino group, N-ethyl-N-isobutylamino  
group, N-(2-ethylbutyl)amino group, N-(2-ethylbutyl)-N-methylamino group,  
N-cyclopentylamino group, N-cyclopentyl-N-methylamino group, N-cyclohexylamino  
group, N-cyclohexyl-N-methylamino group, N-cycloheptylamino group,  
N-(cyclopentylmethyl)amino group, N-(cyclopentylmethyl)-N-methylamino group,  
N-(cyclohexylmethyl)amino group, N-(cyclohexylmethyl)-N-methylamino group,  
N-(2-methylphenyl)amino group, N-(4-methylphenyl)amino group,  
N-(2-fluorophenyl)amino group, N-(3-fluorophenyl)amino group,  
N-(4-fluorophenyl)amino group, N-(2-chlorophenyl)amino group,  
N-(3-chlorophenyl)amino group, N-(4-chlorophenyl)amino group,



N-(indan-2-yl)amino group, N-(1-phenylethyl)amino group,  
 N-[1-(2-fluorophenyl)ethyl]amino group, N-[1-(3-fluorophenyl)ethyl]amino group,  
 N-[1-(4-fluorophenyl)ethyl]amino group, N-[1-(2-chlorophenyl)ethyl]amino group,  
 N-[1-(3-chlorophenyl)ethyl]amino group, N-[1-(4-chlorophenyl)ethyl]amino group,  
 N-(2-methylphenylmethyl)amino group, N-methyl-N-(2-methylphenylmethyl)amino  
 group, N-(3-methylphenylmethyl)amino group,  
 N-methyl-N-(3-methylphenylmethyl)amino group, N-(4-methylphenylmethyl)amino  
 group, N-methyl-N-(4-methylphenylmethyl)amino group,  
 N-(2-fluorophenylmethyl)amino group, N-(2-fluorophenylmethyl)-N-methylamino  
 group, N-(3-fluorophenylmethyl)amino group,  
 N-(3-fluorophenylmethyl)-N-methylamino group, N-(4-fluorophenylmethyl)amino  
 group, N-(4-fluorophenylmethyl)-N-methylamino group,  
 N-(2-chlorophenylmethyl)amino group, N-(2-chlorophenylmethyl)-N-methylamino  
 group, N-(3-chlorophenylmethyl)amino group,  
 N-(3-chlorophenylmethyl)-N-methylamino group, N-(4-chlorophenylmethyl)amino  
 group, N-(4-chlorophenylmethyl)-N-methylamino group,  
 N-(2,3-difluorophenylmethyl)amino group,  
 N-(2,3-difluorophenylmethyl)-N-methylamino group,  
 N-(2,4-difluorophenylmethyl)amino group,  
 N-(2,4-difluorophenylmethyl)-N-methylamino group,  
 N-(2,5-difluorophenylmethyl)amino group,  
 N-(2,5-difluorophenylmethyl)-N-methylamino group,  
 N-(3,4-difluorophenylmethyl)amino group,  
 N-(3,4-difluorophenylmethyl)-N-methylamino group,  
 N-(3,5-difluorophenylmethyl)amino group,  
 N-(3,5-difluorophenylmethyl)-N-methylamino group,  
 N-(2,3-dichlorophenylmethyl)amino group,

N-(2,3-dichlorophenylmethyl)-N-methylamino group,  
 N-(2,4-dichlorophenylmethyl)amino group,  
 N-(2,4-dichlorophenylmethyl)-N-methylamino group,  
 N-(2,5-dichlorophenylmethyl)amino group,  
 N-(2,5-dichlorophenylmethyl)-N-methylamino group,  
 N-(2,6-dichlorophenylmethyl)amino group,  
 N-(2,6-dichlorophenylmethyl)-N-methylamino group,  
 N-(3,4-dichlorophenylmethyl)amino group,  
 N-(3,4-dichlorophenylmethyl)-N-methylamino group,  
 N-(3,5-dichlorophenylmethyl)amino group,  
 N-(3,5-dichlorophenylmethyl)-N-methylamino group,  
 N-[2-(trifluoromethyl)phenylmethyl]amino group,  
 N-methyl-N-[2-(trifluoromethyl)phenylmethyl]amino group,  
 N-[3-(trifluoromethyl)phenylmethyl]amino group,  
 N-methyl-N-[3-(trifluoromethyl)phenylmethyl]amino group,  
 N-[4-(trifluoromethyl)phenylmethyl]amino group,  
 N-methyl-N-[4-(trifluoromethyl)phenylmethyl]amino group, 1-pyrrolidino group,  
 1-(4-methylpiperidino) group, 1-homopiperidino group, or 4-morpholino group,

AR is naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group,  
 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group,  
 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group,  
 6-(N,N-dimethylamino)naphthalen-2-yl group,  
 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group,  
 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group,  
 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group,  
 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group,  
 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group,

2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group,  
 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group,  
 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group,  
 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group,  
 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group,  
 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group,  
 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group,  
 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group,  
 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group,  
 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group,  
 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group,  
 quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl  
 group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group,  
 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group,  
 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group,  
 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group,  
 imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl  
 group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group,  
 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, or benzoxazol-5-yl  
 group, and

Y is hydrogen atom, methyl group, or ethyl group.

In another particularly preferred embodiment of the present invention, the compound represented by the formula (I) or a salt thereof satisfies all of the following requirements.

Link represents  $-(CH_2)_n-$ , symbol n represents an integer of 2.

C<sup>3</sup> represents carbon atom to which AR bonds, C<sup>4</sup> represents carbon atom to which Rs bonds, C<sup>5</sup> represents carbon atom substituted with Zx, and C<sup>2</sup> and C<sup>6</sup> represent unsubstituted ring-constituting carbon atom.

Zx represents N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, and N,N-dimethylsulfamoylamino group.

Rs represents -O-Rx. Rx represents a group as any one of butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group,

2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group,  
 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl  
 group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group,  
 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group,  
 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group,  
 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group,  
 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group,  
 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group,  
 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group,  
 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group,  
 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group,  
 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group,  
 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group,  
 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group,  
 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,  
 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,  
 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group,  
 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group,  
 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group,  
 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group,  
 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group,  
 2-(N-phenyl-N-methylamino)ethyl group, and 2-(N-ethyl-N-phenylamino)ethyl  
 group.

AR represents a group as any one of naphthalen-2-yl group,  
 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group,  
 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group,  
 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl

group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group,  
 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group,  
 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group,  
 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group,  
 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group,  
 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group,  
 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group,  
 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group,  
 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group,  
 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group,  
 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group,  
 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group,  
 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group,  
 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group,  
 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group,  
 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group,  
 quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl  
 group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group,  
 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group,  
 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group,  
 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group,  
 imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group,

1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, and benzoxazol-5-yl group.

The group Y represents hydrogen atom, methyl group, or ethyl group.

In another particularly preferred embodiment of the present invention, the compound represented by the formula (I) or a salt thereof satisfies all of the following requirements.

Link represents  $-(CH_2)_n-$ , symbol n represents an integer of 2.

C<sup>3</sup> represents carbon atom to which AR bonds, C<sup>4</sup> represents carbon atom to which Rs bonds, C<sup>5</sup> may be replaced with V, and C<sup>2</sup> and C<sup>6</sup> represent unsubstituted ring-constituting carbon atom.

V represents nitrogen atom, or carbon atom substituted with Zx, and Zx represents a group as any one of chlorine atom, bromine atom, methyl group, hydroxyl group, methoxy group, amino group, N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, and N,N-dimethylsulfamoylamino group.

R<sub>s</sub> represents -O-R<sub>c</sub>. p in R<sub>c</sub> represents an integer of 2, and A<sup>4</sup> represents a single bond or methylene. A<sup>5</sup> represents -C(O)-, -C(S)-, or -S(O)<sub>2</sub>-. R<sub>d</sub> represents a group as any one of methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, and 4-fluorophenylmethyl group. R<sub>e</sub> represents a group as any one of isopropyl group, butyl group, isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl

group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenyloxy group, 4-methylphenyloxy group, 4-chlorophenyloxy group, 4-fluorophenyloxy group, N-propylamino group, N-isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group, N-(4-fluorophenyl)amino group, pyrrolidino group, piperidino group, and morpholino group.

AR represents a group as any one of naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group,



2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group, 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, and benzoxazol-5-yl group.

The group Y represents hydrogen atom, methyl group, or ethyl group.

In another particularly preferred embodiment of the present invention, the compound represented by the formula (I) or a salt thereof satisfies all of the following requirements.

Link represents  $-(CH_2)_n-$ , symbol n represents an integer of 2.

C<sup>3</sup> represents carbon atom to which AR bonds, C<sup>4</sup> represents carbon atom to which Rs bonds, C<sup>5</sup> may be replaced with V, and C<sup>2</sup> and C<sup>6</sup> represent unsubstituted

ring-constituting carbon atom.

V represents nitrogen atom, or carbon atom substituted with Zx, Zx is any one of fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, or N,N-dimethylamino group,

Rs represents ·D·Rx and D represents a single bond. Rx is butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 2-methylphenyl group, 3-methylphenyl group, 4-methylphenyl group, 2,3-dimethylphenyl group, 3,5-dimethylphenyl group, 2-methoxyphenyl group, 3-methoxyphenyl group, 4-methoxyphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, 2,3-difluorophenyl group, 2,4-difluorophenyl group, 2,5-difluorophenyl group, 3,4-difluorophenyl group, 2,3-dichlorophenyl group, 2,4-dichlorophenyl group, 2,5-dichlorophenyl group, 2,6-dichlorophenyl group, 3,4-dichlorophenyl group, 3,5-dichlorophenyl group, 2-trifluoromethylphenyl group, 3-trifluoromethylphenyl group, 4-trifluoromethylphenyl group, 4-(N,N-dimethylamino)phenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, furan-2-yl group, furan-3-yl group, thiophen-2-yl group, thiophen-3-yl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, naphthalen-1-yl group, naphthalen-2-yl group, 1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, biphenyl-2-yl group, biphenyl 3-yl group,

biphenyl-4-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group,  
 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group,  
 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group,  
 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl  
 group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group,  
 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group,  
 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl  
 group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group,  
 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group,  
 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group,  
 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group,  
 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group,  
 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group,  
 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group,  
 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group,  
 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group,  
 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group,  
 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group,  
 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,  
 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,  
 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group,  
 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group,  
 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group,  
 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group,  
 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group,  
 2-(N-phenyl-N-methylamino)ethyl group, or 2-(N-ethyl-N-phenylamino)ethyl group,

AR is naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group,

6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group,  
 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group,  
 6-(N,N-dimethylamino)naphthalen-2-yl group,  
 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group,  
 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group,  
 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group,  
 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group,  
 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group,  
 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group,  
 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group,  
 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group,  
 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group,  
 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group,  
 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group,  
 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group,  
 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group,  
 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group,  
 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group,  
 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group,  
 quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl  
 group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group,  
 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group,

1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, or benzoxazol-5-yl group, and

Y is hydrogen atom, methyl group, or ethyl group.

In another particularly preferred embodiment of the present invention, the compound represented by the formula (I) or a salt thereof satisfies all of the following requirements.

Link represents  $(CH_2)_n$ , symbol n represents an integer of 2.

C<sup>3</sup> represents carbon atom to which AR bonds, C<sup>4</sup> represents carbon atom to which Rs bonds, C<sup>5</sup> may be replaced with V, and C<sup>2</sup> and C<sup>6</sup> represent unsubstituted ring-constituting carbon atom.

V represents nitrogen atom, or carbon atom substituted with Z<sub>x</sub>, Z<sub>x</sub> is any one of fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, or N,N-dimethylamino group,

Rs represents -D-R<sub>x</sub> and D represents a single bond. R<sub>x</sub> is phenyl group, 2-methylphenyl group, 3-methylphenyl group, 4-methylphenyl group, 2,3-dimethylphenyl group, 3,5-dimethylphenyl group, 2-methoxyphenyl group, 3-methoxyphenyl group, 4-methoxyphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, 2,3-difluorophenyl group, 2,4-difluorophenyl group, 2,5-difluorophenyl group, 3,4-difluorophenyl group, 2,3-dichlorophenyl group, 2,4-dichlorophenyl group, 2,5-dichlorophenyl group, 2,6-dichlorophenyl group,

3,4-dichlorophenyl group, 3,5-dichlorophenyl group, 2-trifluoromethylphenyl group, 3-trifluoromethylphenyl group, 4-trifluoromethylphenyl group, 4-(N,N-dimethylamino)phenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, furan-2-yl group, furan-3-yl group, thiophen-2-yl group, thiophen-3-yl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, naphthalen-1-yl group, naphthalen-2-yl group, 1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1H-indazol-5-yl group, or 1-methyl-1H-indazol-5-yl group,

AR is naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group,

1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group,  
 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group,  
 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group,  
 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group,  
 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group,  
 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group,  
 quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl  
 group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group,  
 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group,  
 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group,  
 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group,  
 imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl  
 group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group,  
 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, or benzoxazol-5-yl  
 group, and

Y is hydrogen atom, methyl group, or ethyl group.

In another particularly preferred embodiment of the present invention, the compound represented by the formula (I) or a salt thereof satisfies all of the following requirements.

Link represents  $-(CH_2)_n-$ , symbol n represents an integer of 2.

C<sup>3</sup> represents carbon atom to which AR bonds, C<sup>4</sup> represents carbon atom to which Rs bonds, and C<sup>2</sup>, C<sup>5</sup>, and C<sup>6</sup> represent unsubstituted ring-constituting carbon atom.

Rs represents -D-Rx and D represents a single bond. Rx is phenyl group, 2-methylphenyl group, 3-methylphenyl group, 4-methylphenyl group, 2,3-dimethylphenyl group, 3,5-dimethylphenyl group, 2-methoxyphenyl group, 3-methoxyphenyl group, 4-methoxyphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, 2,3-difluorophenyl group, 2,4-difluorophenyl group, 2,5-difluorophenyl group, 3,4-difluorophenyl group, 2,3-dichlorophenyl group, 2,4-dichlorophenyl group, 2,5-dichlorophenyl group, 2,6-dichlorophenyl group, 3,4-dichlorophenyl group, 3,5-dichlorophenyl group, 2-trifluoromethylphenyl group, 3-trifluoromethylphenyl group, 4-trifluoromethylphenyl group, 4-(N,N-dimethylamino)phenyl group, indan-2-yl group, furan-2-yl group, furan-3-yl group, thiophen-2-yl group, thiophen-3-yl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, naphthalen-1-yl group, naphthalen-2-yl group, 1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1H-indazol-5-yl group, or 1-methyl-1H-indazol-5-yl group,

AR is naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group,



2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group,  
 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group,  
 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group,  
 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group,  
 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group,  
 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group,  
 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group,  
 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group,  
 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group,  
 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group,  
 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group,  
 quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl  
 group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group,  
 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group,  
 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group,  
 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group,  
 imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl  
 group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group,  
 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, or benzoxazol-5-yl  
 group, and

Y is hydrogen atom, methyl group, or ethyl group.

Compound (I) of the present invention may have one or more asymmetric carbons depending on types of substituents. For example, as for a compound wherein the group Rs contains one or more asymmetric carbons, two kinds of optical isomers exist when the number of asymmetric carbon is 1, and when the number of asymmetric carbons is 2, four kinds of optical isomers and two kinds of diastereomers exist. Pure stereoisomers including optical isomers and diastereoisomers, any mixtures thereof, racemates and the like of the stereoisomers fall within the scope of the present invention. Further, Compound (I) of the present invention may exist as geometrical isomers based on a cycloalkyl ring structure, and any geometrical isomers in pure forms, and any mixtures of the geometrical isomers also fall within the scope of the present invention. Mixtures such as racemates may sometimes be preferred from a viewpoint of easiness for manufacture.

As a salt of Compound (I) of the present invention, a pharmaceutically acceptable salt is preferred. It is meant that, when at least one of the conditions (1) to (3) is satisfied: (1) Y is hydrogen atom; (2) the group AR contains carboxyl group or phenolic hydroxyl group; (3) the group Zx is phenolic hydroxyl group, and the like, then the compound forms 1 to 3 alkali salts depending on the number of acidic groups. Examples the alkali salts include, for example, salts with inorganic bases such as sodium and ammonia and salts with organic bases such as triethylamine.

Alternatively, it is meant that, when at least one of the conditions (1) to (4) is satisfied: (1) the group Rs has properties as a base as in a compound wherein Rs contains a substituted or unsubstituted amino group and the like; (2) AR itself is a cyclic substituent having properties as a base; (3) the group Ar contains a substituted or unsubstituted amino group; (4) any carbon atom in the aromatic ring (E) is replaced with V, and V is nitrogen atom, V is carbon atom substituted with Zx, and Zx is a substituted or unsubstituted amino group and the like, then the

compound forms 1 to 4 acidic salts depending on the number of basic groups.

Examples of the acidic salts include, for example, salts with inorganic acids such as hydrochloric acid and sulfuric acid and salts with organic acids such as acetic acid and citric acid.

C<sup>2'</sup>, C<sup>3'</sup>, C<sup>4'</sup>, C<sup>5'</sup>, and C<sup>6'</sup> in the aromatic ring (E') in the aforementioned formula (II) each represent a ring-constituting carbon atom. Among them, any ring-constituting carbon atom to which Rs' and G do not bind may be replaced with V'. The substitution positions of Rs', G, and V' are similar to those described in the explanations of the substitution positions of Rs (corresponding to the position of Rs'), AR (corresponding to the position of the group G), and V (corresponding to the position of V') in the aforementioned formula (I).

V' represents nitrogen atom, or represents carbon atom substituted with Zx'. Zx' has the same meaning as that of Zx, provided that when Zx contains hydroxyl group (OH), the hydroxyl group may be protected with Rp<sup>1</sup>, and when Zx contains amino group (NH), the amino group may be protected with Rp<sup>2</sup>.

Rs' represents -D·Rx' or -N(Ry')(Rz'). -D·Rx' and -N(Ry')(Rz') have the same meanings as those of -D·Rx and -N(Ry)(Rz) mentioned above, respectively. Provided that when -D·Rx and -N(Ry)(Rz) contain hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when -D·Rx and -N(Ry)(Rz) contains amino group (NH), the amino group may be protected with Rp<sup>2</sup>.

Rp<sup>1</sup> represents, for example, a silyl group substituted with 3 of identical or different linear or branched saturated alkyl groups having 1 to 4 carbon atoms or phenyl groups, tetrahydropyranyl group, tetrahydrofuryl group, allyl group, propargyl group, benzyl group which may be substituted with one T<sup>1</sup> or two or more identical or different T<sup>1</sup>, -CH<sub>2</sub>-U·Rp<sup>3</sup>, -C(O)Rp<sup>3</sup>, -C(O)ORp<sup>3</sup>, or the like. U represents oxygen atom, or sulfur atom, and Rp<sup>3</sup> represents hydrogen atom, a linear or branched saturated alkyl group having 1 to 4 carbon atoms, trimethylsilylethyl

group, chloromethyl group, trichloromethyl group, trifluoromethyl group, 9-fluorenylmethyl group, adamantyl group, allyl group, -A<sup>6</sup>-Qp, or the like. Rp<sup>2</sup> represents, for example, benzyl group which may be substituted with one of T<sup>1</sup> or two or more of identical or different T<sup>1</sup>, -C(O)Rp<sup>3</sup>, -C(O)ORp<sup>3</sup>, or the like. However, the protective groups of hydroxyl group and amino group are not limited to these, and they can be chosen by referring and examining methods for introduction of protective groups and deprotection described in usual publications in the chemical field, for example, Protective Groups In Organic Synthesis, THIRD EDITION, published by John Wiley & Sons or the references cited therein.

G represents chlorine atom, bromine atom, iodine atom, mesylate group, triflate group, or an arenesulfonate group of which aromatic moiety may be substituted with one of T<sup>1</sup> or two or more identical or different T<sup>1</sup>. Examples of the arenesulfonate group include, for example, benzenesulfonate group, p-toluenesulfonate group, mesitylenesulfonate group, 2,4,6-triisopropylbenzenesulfonate group, 4-fluorobenzenesulfonate group, 2,5-dichlorobenzenesulfonate group, 3-(trifluoromethyl)benzenesulfonate group, pentafluorobenzenesulfonate group, 2-nitrobenzenesulfonate group, 2,4-dinitrobenzenesulfonate group, and the like. Preferred examples of G include chlorine atom, bromine atom, iodine atom, triflate group, and the like, and bromine atom and iodine atom are particularly preferred examples.

Y' represents a lower alkyl group having 1 to 4 carbon atoms. Examples of the lower alkyl group having 1 to 4 carbon atoms include methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, and the like. Among these, methyl group, and ethyl group are particularly preferred examples.

In the aforementioned formula (II), n and D have the same meaning as defined above.

In a preferred embodiment, the compound represented by the formula (II) satisfies all of the following requirements.

Symbol  $n$  represents an integer of 1 to 3.

The group  $G$  binds to  $C^{2'}$ ,  $Rs'$  binds to any of the atoms  $C^{3'}$ ,  $C^{4'}$  and  $C^{5'}$ , and a ring-constituting carbon atom to which  $Rs'$  does not bind among  $C^{3'}$ ,  $C^{4'}$ , and  $C^{5'}$  may be substituted with  $V'$ .

$V'$  represents nitrogen atom, or carbon atom substituted with  $Zx'$ , and  $Zx'$  represents any one of fluorine atom, chlorine atom, bromine atom, nitro group, methyl group, hydroxyl group, methoxy group, amino group,  $N$ -methylamino group,  $N$ -ethylamino group,  $N$ -propylamino group,  $N$ -isopropylamino group,  $N,N$ -dimethylamino group,  $N,N$ -diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, and  $N,N$ -dimethylsulfamoylamino group, provided that when  $Zx'$  contains hydroxyl group, the hydroxyl group may be protected with  $Rp^1$ , and when the substituted  $Zx'$  contains amino group, the amino group may be protected with  $Rp^2$ .

$Rs'$  represents  $-D-Rx'$  or  $-N(Ry')(Rz')$ .  $D$  represents oxygen atom or sulfur atom.  $Rx'$  represents butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-cyclopentylethyl group, or 2-cyclohexylethyl group, or represents  $Rb$  or  $Rc$ .  $Q$  in  $Rb$  represents a group as any one of phenyl group, thienyl group, furyl group, pyridyl group, oxazolyl group, naphthyl group, tetrahydronaphthyl group, indanyl group, indolyl group, and dihydrobenzodioxyl group.  $A^2$  represents a single bond, oxygen atom, sulfur atom,  $-N(methyl)-$ , or  $-N(ethyl)-$  (provided that when  $A^2$  represents oxygen atom, sulfur atom,  $-N(methyl)-$ , or  $-N(ethyl)-$ ,  $A^1$  represents ethylene).  $R^2$  and  $R^3$  independently represent hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, dimethylamino group, acetylamino group, or methylsulfonylamino group

(provided that when Q represents phenyl group, A<sup>1</sup> represents a single bond, or unsubstituted methylene, and A<sup>2</sup> represents a single bond, one of R<sup>2</sup> and R<sup>3</sup> represents a substituent other than hydrogen atom). Symbol p in R<sub>c</sub> represents an integer of 2 or 3, and A<sup>4</sup> represents a single bond or methylene. A<sup>5</sup> represents -C(O)-, -C(S)-, or -S(O)<sub>2</sub>-. R<sub>d</sub> represents hydrogen atom, or a group as any one of methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group, cyclopropylmethyl group, cyclopentyl group, cyclopentylmethyl group, cyclohexyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, and pyridin-4-yl group. R<sub>e</sub> represents any one of methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, phenylmethyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, furan-2-yl group, furan-3-yl group, thiophen-2-yl group, thiophen-3-yl group, methoxy group, ethoxy group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenyloxy group, 4-methylphenyloxy group, 4-chlorophenyloxy group, 4-fluorophenyloxy group, thiomethoxy group, amino group, N-methylamino group, N,N-dimethylamino group, N-ethylamino group, N,N-diethylamino group, N-propylamino group, N-isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group,

N-(4-fluorophenyl)amino group, N-(pyridin-2-yl)amino group, N-(pyridin-3-yl)amino group, N-(pyridin-4-yl)amino group, N-(furan-2-yl)amino group, N-(furan-3-yl)amino group, N-(thiophen-2-yl)amino group, N-(thiophen-3-yl)amino group, pyrrolidino group, piperidino group, morpholino group, methyloxycarbonylamino group, and ethyloxycarbonylamino group. Rz' represents any one of butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group,

3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group,  
 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group,  
 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group,  
 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group,  
 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group,  
 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group,  
 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,  
 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,  
 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group,  
 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group,  
 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group,  
 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group,  
 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group,  
 2-(N-phenyl-N-methylamino)ethyl group, 2-(N-ethyl-N-phenylamino)ethyl group,  
 isobutyryl group, isopropylthiocarbonyl group, isopropylsulfonyl group, valeryl  
 group, butylthiocarbonyl group, isovaleryl group, isobutylthiocarbonyl group,  
 pivaloyl group, t-butylthiocarbonyl group, cyclopropylcarbonyl group,  
 cyclopropylthiocarbonyl group, cyclopentylcarbonyl group, cyclopentylthiocarbonyl  
 group, cyclohexylcarbonyl group, cyclohexylthiocarbonyl group,  
 cyclopentylmethylcarbonyl group, cyclopentylmethylthiocarbonyl group,  
 cyclohexylmethylcarbonyl group, cyclohexylmethylthiocarbonyl group, benzoyl  
 group, thiobenzoyl group, phenylsulfonyl group, 4-methylphenylcarbonyl group,  
 4-methylphenylthiocarbonyl group, 4-methylphenylsulfonyl group,  
 4-chlorophenylcarbonyl group, 4-chlorophenylthiocarbonyl group,  
 4-fluorophenylcarbonyl group, 4-fluorophenylthiocarbonyl group,  
 isopropylloxycarbonyl group, N-isopropylcarbamoyl group, N-isopropylthiocarbamoyl  
 group, butylloxycarbonyl group, N-butylcarbamoyl group, N-butylthiocarbamoyl



group, isobutyloxycarbonyl group, N-isobutylcarbamoyl group, N-isobutylthiocarbamoyl group, t-butyloxycarbonyl group, N-t-butylcarbamoyl group, N-t-butylthiocarbamoyl group, cyclopropyloxycarbonyl group, N-cyclopropylcarbamoyl group, N-cyclopropylthiocarbamoyl group, cyclopentyloxycarbonyl group, N-cyclopentylcarbamoyl group, N-cyclopentylthiocarbamoyl group, cyclohexyloxycarbonyl group, N-cyclohexylcarbamoyl group, N-cyclohexylthiocarbamoyl group, cyclopentylmethyloxycarbonyl group, cyclohexylmethyloxycarbonyl group, phenyloxycarbonyl group, N-phenylcarbamoyl group, N-phenylthiocarbamoyl group, 4-methylphenyloxycarbonyl group, N-(4-methylphenyl)carbamoyl group, N-(4-methylphenyl)thiocarbamoyl group, 4-chlorophenyloxycarbonyl group, N-(4-chlorophenyl)carbamoyl group, N-(4-chlorophenyl)thiocarbamoyl group, 4-fluorophenyloxycarbonyl group, N-(4-fluorophenyl)carbamoyl group, N-(4-fluorophenyl)thiocarbamoyl group, (pyrrolidino-1-yl)carbonyl group, (piperidino-1-yl)carbonyl group, and (morpholino-4-yl)carbonyl group. Ry' represents hydrogen atom, methyl group, ethyl group, or isobutyl group, or binds to Rz' to form pyrrolidino group, piperidino group, piperazino group, morpholino group, pyrrol-1-yl group, imidazol-1-yl group, or pyrazol-1-yl group together with the nitrogen atom to which they bonds. Provided that when -D-Rx' or -N(Ry')(Rz') contains hydroxyl group (OH), the hydroxyl group may be protected with Rp<sup>1</sup>, and when -D-Rx' or -N(Ry')(Rz') contains amino group, the amino group may be protected with Rp<sup>2</sup>.

The group G represents chlorine atom, bromine atom, iodine atom, or triflate group.

The group Y' represents methyl group, or ethyl group.

In another preferred embodiment, the compound represented by the formula (II) satisfies all of the following requirements.

Symbol  $n$  represents an integer of 1 to 3.

The group  $G$  binds to  $C^3$ ,  $Rs'$  binds to any of the atoms  $C^4$ ,  $C^5$ , and  $C^6$ , and a ring-constituting carbon atom to which  $Rs'$  does not bind among  $C^4$ ,  $C^5$  and  $C^6$  may be replaced with  $V'$ .

$V'$  represents nitrogen atom, or carbon atom substituted with  $Zx'$ , and  $Zx'$  represents any one of fluorine atom, chlorine atom, bromine atom, nitro group, methyl group, hydroxyl group, methoxy group, amino group,  $N$ -methylamino group,  $N$ -ethylamino group,  $N$ -propylamino group,  $N$ -isopropylamino group,  $N,N$ -dimethylamino group,  $N,N$ -diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, and  $N,N$ -dimethylsulfamoylamino group, provided that when  $Zx'$  contains hydroxyl group, the hydroxyl group may be protected with  $Rp^1$ , and when the substituted  $Zx'$  contains amino group, the amino group may be protected with  $Rp^2$ .

$Rs'$  represents  $-D-Rx'$ , or  $-N(Ry')(Rz')$ .  $D$  represents oxygen atom or sulfur atom.  $Rx'$  represents butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-cyclopentylethyl group, or 2-cyclohexylethyl group, or represents  $Rb$  or  $Rc$ .  $Q$  in  $Rb$  represents a group as any one of phenyl group, thienyl group, furyl group, pyridyl group, oxazolyl group, naphthyl group, tetrahydronaphthyl group, indanyl group, indolyl group, and dihydrobenzodioxyl group.  $A^2$  represents a single bond, oxygen atom, sulfur atom,  $-N(methyl)-$ , or  $-N(ethyl)-$  (provided that when  $A^2$  represents oxygen atom, sulfur atom,  $-N(methyl)-$ , or  $-N(ethyl)-$ ,  $A^1$  represents ethylene).  $R^2$  and  $R^3$  independently represent hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, dimethylamino group, acetylamino group, or methylsulfonylamino group (provided that when  $Q$  represents phenyl group,  $A^1$  represents a single bond, or unsubstituted methylene, and  $A^2$  represents a single bond, one of  $R^2$  and  $R^3$

represents a substituent other than hydrogen atom). Symbol p in R<sub>c</sub> represents an integer of 2 or 3, and A<sup>4</sup> represents a single bond or methylene. A<sup>5</sup> represents -C(O)-, -C(S)-, or -S(O)<sub>2</sub>-. R<sub>d</sub> represents hydrogen atom, or a group as any one of methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group, cyclopropylmethyl group, cyclopentyl group, cyclopentylmethyl group, cyclohexyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, and pyridin-4-yl group. R<sub>e</sub> represents any one of methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, phenylmethyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, furan-2-yl group, furan-3-yl group, thiophen-2-yl group, thiophen-3-yl group, methoxy group, ethoxy group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenyloxy group, 4-methylphenyloxy group, 4-chlorophenyloxy group, 4-fluorophenyloxy group, thiomethoxy group, amino group, N-methylamino group, N,N-dimethylamino group, N-ethylamino group, N,N-diethylamino group, N-propylamino group, N-isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group, N-(4-fluorophenyl)amino group, N-(pyridin-2-yl)amino group, N-(pyridin-3-yl)amino group, N-(pyridin-4-yl)amino group, N-(furan-2-yl)amino group, N-(furan-3-yl)amino

group, N-(thiophen-2-yl)amino group, N-(thiophen-3-yl)amino group, pyrrolidino group, piperidino group, morpholino group, methyloxycarbonylamino group, and ethyloxycarbonylamino group. R<sub>2</sub>' represents any one of butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group,

4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group,  
 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group,  
 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group,  
 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group,  
 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,  
 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,  
 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group,  
 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group,  
 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group,  
 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group,  
 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group,  
 2-(N-phenyl-N-methylamino)ethyl group, 2-(N-ethyl-N-phenylamino)ethyl group,  
 isobutyryl group, isopropylthiocarbonyl group, isopropylsulfonyl group, valeryl  
 group, butylthiocarbonyl group, isovaleryl group, isobutylthiocarbonyl group,  
 pivaloyl group, t-butylthiocarbonyl group, cyclopropylcarbonyl group,  
 cyclopropylthiocarbonyl group, cyclopentylcarbonyl group, cyclopentylthiocarbonyl  
 group, cyclohexylcarbonyl group, cyclohexylthiocarbonyl group,  
 cyclopentylmethylcarbonyl group, cyclopentylmethylthiocarbonyl group,  
 cyclohexylmethylcarbonyl group, cyclohexylmethylthiocarbonyl group, benzoyl  
 group, thiobenzoyl group, phenylsulfonyl group, 4-methylphenylcarbonyl group,  
 4-methylphenylthiocarbonyl group, 4-methylphenylsulfonyl group,  
 4-chlorophenylcarbonyl group, 4-chlorophenylthiocarbonyl group,  
 4-fluorophenylcarbonyl group, 4-fluorophenylthiocarbonyl group,  
 isopropylloxycarbonyl group, N-isopropylcarbamoyl group, N-isopropylthiocarbamoyl  
 group, butyloxycarbonyl group, N-butylcarbamoyl group, N-butylthiocarbamoyl  
 group, isobutyloxycarbonyl group, N-isobutylcarbamoyl group,  
 N-isobutylthiocarbamoyl group, t-butyloxycarbonyl group, N-t-butylcarbamoyl

group, N-t-butylthiocarbamoyl group, cyclopropyloxycarbonyl group, N-cyclopropylcarbamoyl group, N-cyclopropylthiocarbamoyl group, cyclopentyloxycarbonyl group, N-cyclopentylcarbamoyl group, N-cyclopentylthiocarbamoyl group, cyclohexyloxycarbonyl group, N-cyclohexylcarbamoyl group, N-cyclohexylthiocarbamoyl group, cyclopentylmethyloxycarbonyl group, cyclohexylmethyloxycarbonyl group, phenyloxycarbonyl group, N-phenylcarbamoyl group, N-phenylthiocarbamoyl group, 4-methylphenyloxycarbonyl group, N-(4-methylphenyl)carbamoyl group, N-(4-methylphenyl)thiocarbamoyl group, 4-chlorophenyloxycarbonyl group, N-(4-chlorophenyl)carbamoyl group, N-(4-chlorophenyl)thiocarbamoyl group, 4-fluorophenyloxycarbonyl group, N-(4-fluorophenyl)carbamoyl group, N-(4-fluorophenyl)thiocarbamoyl group, (pyrrolidino-1-yl)carbonyl group, (piperidino-1-yl)carbonyl group, and (morpholino-4-yl)carbonyl group. Ry' represents hydrogen atom, methyl group, ethyl group, or isobutyl group, or binds to Rz' to form pyrrolidino group, piperidino group, piperazino group, morpholino group, pyrrol-1-yl group, imidazol-1-yl group, or pyrazol-1-yl group together with nitrogen atom. Provided that when -D-Rx' or -N(Ry')(Rz') contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and -D-Rx' or -N(Ry')(Rz') contains amino group, the amino group may be protected with Rp<sup>2</sup>.

The group G represents chlorine atom, bromine atom, iodine atom, or triflate group.

The group Y' represents methyl group, or ethyl group.

In a particularly preferred embodiment, the compound represented by the formula (II) satisfies all of the following requirements.

Symbol n represents an integer of 2.

C<sup>2'</sup> represents carbon atom to which the group G bonds, C<sup>3'</sup> represents carbon atom to which Rs' binds, C<sup>4'</sup> may be replaced with V', and C<sup>5'</sup> and C<sup>6'</sup> represent

an unsubstituted ring-constituting carbon atom.

V' represents nitrogen atom, or carbon atom substituted with Zx', and Zx' represents any one of fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, and N,N-dimethylamino group, provided that when Zx' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when the substituted Zx' contains amino group, the amino group may be protected with Rp<sup>2</sup>.

Rs' represents -O-Rx'. Rx' represents any one of butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group,

3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group,  
 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group,  
 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group,  
 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group,  
 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group,  
 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group,  
 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group,  
 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group,  
 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group,  
 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,  
 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,  
 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group,  
 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group,  
 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group,  
 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group,  
 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group,  
 2-(N-phenyl-N-methylamino)ethyl group, and 2-(N-ethyl-N-phenylamino)ethyl  
 group.

The group G represents bromine atom, or iodine atom.

The group Y' represents methyl group, or ethyl group.

In another particularly preferred embodiment, the compound represented by the formula (II) satisfies all of the following requirements.

Symbol n represents an integer of 2.

C<sup>2'</sup> represents carbon atom to which the group G bonds, C<sup>4'</sup> represents carbon atom to which Rs' binds, C<sup>5'</sup> may be replaced with V', and C<sup>3'</sup> and C<sup>6'</sup> represent an unsubstituted ring-constituting carbon atom.

V' represents nitrogen atom, or carbon atom substituted with Zx', and Zx'



represents any one of fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, and N,N-dimethylamino group, provided that when Zx' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when the substituted Zx' contains amino group, the amino group may be protected with Rp<sup>2</sup>.

Rs' represents -O-Rx'. Rx' represents any one of butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group,

2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group,  
 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group,  
 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group,  
 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group,  
 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group,  
 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group,  
 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group,  
 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,  
 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,  
 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group,  
 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group,  
 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group,  
 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group,  
 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group,  
 2-(N-phenyl-N-methylamino)ethyl group, and 2-(N-ethyl-N-phenylamino)ethyl  
 group.

The group G represents bromine atom, or iodine atom.

The group Y' represents methyl group, or ethyl group.

In another particularly preferred embodiment, the compound represented by the formula (II) satisfies all of the following requirements.

Symbol n represents an integer of 2.

C<sup>3'</sup> represents carbon atom to which the group G bonds, C<sup>5'</sup> represents carbon atom to which Rs' binds, and C<sup>2'</sup>, C<sup>4'</sup> and C<sup>6'</sup> represent an unsubstituted ring-constituting carbon atom.

R<sub>s</sub>' represents -O-R<sub>x</sub>'. R<sub>x</sub>' represents any one of butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group,

4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,

2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,  
 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group,  
 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group,  
 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group,  
 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group,  
 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group,  
 2-(N-phenyl-N-methylamino)ethyl group, and 2-(N-ethyl-N-phenylamino)ethyl  
 group.

The group G represents bromine atom, or iodine atom.

The group Y' represents methyl group, or ethyl group.

In another particularly preferred embodiment, the compound represented by the formula (II) satisfies all of the following requirements.

Symbol n represents an integer of 2.

C<sup>3'</sup> represents carbon atom to which the group G bonds, C<sup>4'</sup> represents carbon atom to which Rs' binds, C<sup>5'</sup> represents nitrogen atom, and C<sup>2'</sup> and C<sup>6'</sup> represent an unsubstituted ring-constituting carbon atom.

Rs' represents -O-Rx'. Rx' represents any one of butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group,

1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group,  
 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group,  
 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl  
 group, 3-methylphenylmethyl group, 4-methylphenylmethyl group,  
 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group,  
 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl  
 group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group,  
 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group,  
 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group,  
 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group,  
 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group,  
 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group,  
 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group,  
 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group,  
 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group,  
 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group,  
 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group,  
 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group,  
 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,  
 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,  
 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group,  
 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group,  
 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group,  
 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group,  
 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group,  
 2-(N-phenyl-N-methylamino)ethyl group, and 2-(N-ethyl-N-phenylamino)ethyl  
 group.

The group G represents bromine atom, or iodine atom.

The group Y' represents methyl group, or ethyl group.

In another particularly preferred embodiment, the compound represented by the formula (II) satisfies all of the following requirements.

Symbol n represents an integer of 2.

C<sup>3'</sup> represents carbon atom to which the group G bonds, C<sup>4'</sup> represents carbon atom to which Rs' binds, C<sup>6'</sup> represents carbon atom substituted with Zx', and C<sup>2'</sup> and C<sup>5'</sup> represent an unsubstituted ring-constituting carbon atom.

Zx' represents any one of fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, and N,N-dimethylamino group, provided that when Zx' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when the substituted Zx' contains amino group, the amino group may be protected with Rp<sup>2</sup>.

Rs' represents -O-Rx'. Rx' represents any one of butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl

group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, and 2-(N-ethyl-N-phenylamino)ethyl group.

The group G represents bromine atom, or iodine atom.

The group Y' represents methyl group, or ethyl group.

In another particularly preferred embodiment, the compound represented by

the formula (II) satisfies all of the following requirements.

Symbol  $n$  represents an integer of 2.

$C^3$  represents carbon atom to which the group  $G$  bonds,  $C^4$  represents carbon atom to which  $Rs'$  binds,  $C^5$  represents carbon atom substituted with  $Zx'$ , or unsubstituted carbon atom, and  $C^2$  and  $C^6$  represent an unsubstituted ring-constituting carbon atom.

$Zx'$  represents any one of fluorine atom, methyl group, hydroxyl group, amino group,  $N$ -methylamino group, and  $N,N$ -dimethylamino group, provided that when  $Zx'$  contains hydroxyl group, the hydroxyl group may be protected with  $Rp^1$ , and when the substituted  $Zx'$  contains amino group, the amino group may be protected with  $Rp^2$ .

$Rs'$  represents  $-S-Rx'$ .  $Rx'$  represents any one of butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group,



2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group, 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group, 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group, 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group, 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group, 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group, 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, and 2-(N-ethyl-N-phenylamino)ethyl group.

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C<sup>3'</sup> represents carbon atom to which the group G bonds, C<sup>4'</sup> represents carbon atom to which Rs' binds, C<sup>5'</sup> represents carbon atom substituted with Zx', or unsubstituted ring-constituting carbon atom, and C<sup>2'</sup> and C<sup>6'</sup> represent an unsubstituted ring-constituting carbon atom.

Zx' represents any one of fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, and N,N-dimethylamino group, provided that when Zx' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when the substituted Zx' contains amino group, the amino group may be protected with Rp<sup>2</sup>.

Rs' represents -N(Ry')(Rz'). Rz' represents any one of butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group,

4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group,  
 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group,  
 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group,  
 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group,  
 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group,  
 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group,  
 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group,  
 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group,  
 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group,  
 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group,  
 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group,  
 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,  
 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,  
 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group,  
 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group,  
 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group,  
 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group,  
 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group,  
 2-(N-phenyl-N-methylamino)ethyl group, 2-(N-ethyl-N-phenylamino)ethyl group,  
 isobutyryl group, isopropylthiocarbonyl group, isopropylsulfonyl group, valeryl  
 group, butylthiocarbonyl group, isovaleryl group, isobutylthiocarbonyl group,  
 pivaloyl group, t-butylthiocarbonyl group, cyclopropylcarbonyl group,  
 cyclopropylthiocarbonyl group, cyclopentylcarbonyl group, cyclopentylthiocarbonyl  
 group, cyclohexylcarbonyl group, cyclohexylthiocarbonyl group,  
 cyclopentylmethylcarbonyl group, cyclopentylmethylthiocarbonyl group,  
 cyclohexylmethylcarbonyl group, cyclohexylmethylthiocarbonyl group, benzoyl  
 group, thiobenzoyl group, phenylsulfonyl group, 4-methylphenylcarbonyl group,

4-methylphenylthiocarbonyl group, 4-methylphenylsulfonyl group, 4-chlorophenylcarbonyl group, 4-chlorophenylthiocarbonyl group, 4-fluorophenylcarbonyl group, 4-fluorophenylthiocarbonyl group, isopropoxy carbonyl group, N-isopropyl carbamoyl group, N-isopropylthiocarbamoyl group, butyloxy carbonyl group, N-butyl carbamoyl group, N-butylthiocarbamoyl group, isobutyloxy carbonyl group, N-isobutyl carbamoyl group, N-isobutylthiocarbamoyl group, t-butyloxy carbonyl group, N-t-butyl carbamoyl group, N-t-butylthiocarbamoyl group, cyclopropyloxy carbonyl group, N-cyclopropyl carbamoyl group, N-cyclopropylthiocarbamoyl group, cyclopentyloxy carbonyl group, N-cyclopentyl carbamoyl group, N-cyclopentylthiocarbamoyl group, cyclohexyloxy carbonyl group, N-cyclohexyl carbamoyl group, N-cyclohexylthiocarbamoyl group, cyclopentylmethyloxy carbonyl group, cyclohexylmethyloxy carbonyl group, phenyloxy carbonyl group, N-phenyl carbamoyl group, N-phenylthiocarbamoyl group, 4-methylphenyloxy carbonyl group, N-(4-methylphenyl) carbamoyl group, N-(4-methylphenyl)thiocarbamoyl group, 4-chlorophenyloxy carbonyl group, N-(4-chlorophenyl) carbamoyl group, N-(4-chlorophenyl)thiocarbamoyl group, 4-fluorophenyloxy carbonyl group, N-(4-fluorophenyl) carbamoyl group, N-(4-fluorophenyl)thiocarbamoyl group, (pyrrolidino-1-yl) carbonyl group, (piperidino-1-yl) carbonyl group, and (morpholino-4-yl) carbonyl group. Ry' represents hydrogen atom, methyl group, ethyl group, or isobutyl group, or binds to Rz' to form pyrrolidino group, piperidino group, or morpholino group together with the nitrogen atom to which they bonds. Provided that when N(Ry')(Rz') contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when Ry' or Rz' contains amino group, the amino group may be protected with Rp<sup>2</sup>.

The group G represents bromine atom, or iodine atom.

The group Y' represents methyl group, or ethyl group.

In another particularly preferred embodiment, the compound represented by the formula (II) satisfies all of the following requirements.

Symbol  $n$  represents an integer of 2.

$C^3$  represents carbon atom to which the group  $G$  bonds,  $C^4$  represents carbon atom to which  $Rs'$  binds,  $C^5$  represents carbon atom substituted with  $Zx'$ , and  $C^2$  and  $C^6$  represent an unsubstituted ring-constituting carbon atom.

$Zx'$  represents any one of N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, and N,N-dimethylsulfamoylamino group. Provided that when the substituted  $Zx'$  contains amino group (NH), the amino group may be protected with  $Rp^2$ .

$Rs'$  represents  $-O \cdot Rx'$ .  $Rx'$  represents any one of butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group,

2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group,  
 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl  
 group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group,  
 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group,  
 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group,  
 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group,  
 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group,  
 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group,  
 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group,  
 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group,  
 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group,  
 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group,  
 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group,  
 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group,  
 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,  
 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,  
 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group,  
 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group,  
 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group,  
 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group,  
 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group,  
 2-(N-phenyl-N-methylamino)ethyl group, and 2-(N-ethyl-N-phenylamino)ethyl  
 group.

The group G represents bromine atom, or iodine atom.

The group Y' represents methyl group, or ethyl group.

In another particularly preferred embodiment, the compound represented by the formula (II) satisfies all of the following requirements.

Symbol  $n$  represents an integer of 2.

$C^3$  represents carbon atom to which the group  $G$  bonds,  $C^4$  represents carbon atom to which  $Rs'$  binds,  $C^5$  represents carbon atom substituted with  $Zx'$ , or unsubstituted carbon atom, and  $C^2$  and  $C^6$  represent an unsubstituted ring-constituting carbon atom.

$Zx'$  represents any one of fluorine atom, methyl group, hydroxyl group, amino group,  $N$ -methylamino group, and  $N,N$ -dimethylamino group, provided that when  $Zx'$  contains hydroxyl group, the hydroxyl group may be protected with  $Rp^1$ , and when the substituted  $Zx'$  contains amino group, the amino group may be protected with  $Rp^2$ .

$Rs'$  represents  $-O-Rx'$ .  $Rx'$  have the same meaning as that of  $Rc$ , provided that when  $Rc$  contains hydroxyl group ( $OH$ ), the hydroxyl group may be protected with  $Rp^1$ .  $p$  in  $Rc$  represents an integer of 2, and  $A^4$  represents a single bond or methylene.  $A^5$  represents  $-C(O)-$ ,  $-C(S)-$ , or  $-S(O)_2-$ .  $Rd$  represents a group as any one of methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, and 4-fluorophenylmethyl group.  $Re$  represents a group as any one of isopropyl group, butyl group, isobutyl group,  $t$ -butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group,  $t$ -butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenyloxy group, 4-methylphenyloxy group, 4-chlorophenyloxy group, 4-fluorophenyloxy group,  $N$ -propylamino group,  $N$ -isopropylamino group,  $N$ -butylamino group,  $N$ -isobutylamino group,  $N$ - $t$ -butylamino group,

N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group, N-(4-fluorophenyl)amino group, pyrrolidino group, piperidino group, and morpholino group.

The group G represents bromine atom, or iodine atom.

The group Y represents methyl group, or ethyl group.

C<sup>2'</sup>, C<sup>3'</sup>, C<sup>4'</sup>, C<sup>5'</sup>, and C<sup>6'</sup> in the aromatic ring (E') in the aforementioned formula (III) each represent a ring-constituting carbon atom. Any ring-constituting carbon atom to which Rs' and AR' do not bond among them may be replaced with V'. The substitution positions of Rs', AR', and V' are similar to those described in the explanations of the substitution positions of Rs (corresponding to the position of Rs'), AR (corresponding to the position of the group AR'), and V (corresponding to the position of V') in the aforementioned formula (I).

AR' has the same meaning as that of AR mentioned above, provided that when AR contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>. In this case, the hydroxyl group includes OH in carboxyl group (COOH). When the substituted AR contains amino group, the amino group represents a substituent, which may be protected with Rp<sup>2</sup>. Examples of the amino group, which may be protected include NH present in a ring constituting AR, for example, as in indole ring, indazole ring, and the like.

Rs', V', n, and D in the aforementioned formula (III) have the same meanings as those defined above. Rp<sup>1</sup>, and Rp<sup>2</sup> also have the same meanings as those defined above.

In a preferred embodiment, the compound represented by the formula (III) satisfies all of the following requirements.

AR' binds to C<sup>2'</sup>, Rs' binds to any of the atoms C<sup>3'</sup>, C<sup>4'</sup>, and C<sup>5'</sup>, and a ring-constituting carbon atom to which Rs' does not bind among C<sup>3'</sup>, C<sup>4'</sup>, and C<sup>5'</sup> may



be replaced with V'.

V' represents nitrogen atom, or carbon atom substituted with Zx', and Zx' represents any one of fluorine atom, chlorine atom, bromine atom, nitro group, methyl group, hydroxyl group, methoxy group, amino group, N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, and N,N-dimethylsulfamoylamino group, provided that when Zx' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when the substituted Zx' contains amino group, the amino group may be protected with Rp<sup>2</sup>.

Rs' represents -D-Rx' or -N(Ry')(Rz'). D represents oxygen atom or sulfur atom. Rx' represents butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-cyclopentylethyl group, or 2-cyclohexylethyl group, or represents Rb or Rc. Q in Rb represents a group as any one of phenyl group, thienyl group, furyl group, pyridyl group, oxazolyl group, naphthyl group, tetrahydronaphthyl group, indanyl group, indolyl group, and dihydrobenzodioxyl group. A<sup>2</sup> represents a single bond, oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)- (provided that when A<sup>2</sup> represents oxygen atom, sulfur atom, -N(methyl)-, or -N(ethyl)-, A<sup>1</sup> represents ethylene). R<sup>2</sup> and R<sup>3</sup> independently represent hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, dimethylamino group, acetylamino group, or methylsulfonylamino group (provided that when Q represents phenyl group, A<sup>1</sup> represents a single bond, or unsubstituted methylene, and A<sup>2</sup> represents a single bond, one of R<sup>2</sup> and R<sup>3</sup> represents a substituent other than hydrogen atom). Symbol p in Rc represents an integer of 2 or 3, and A<sup>4</sup> represents a single bond or methylene. A<sup>5</sup> represents -C(O)-, -C(S)-, or -S(O)<sub>2</sub>-. Rd represents hydrogen atom, or a group as any one of

methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group, cyclopropylmethyl group, cyclopentyl group, cyclopentylmethyl group, cyclohexyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, and pyridin-4-yl group. Re represents any one of methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, phenylmethyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, furan-2-yl group, furan-3-yl group, thiophen-2-yl group, thiophen-3-yl group, methoxy group, ethoxy group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenyloxy group, 4-methylphenyloxy group, 4-chlorophenyloxy group, 4-fluorophenyloxy group, thiomethoxy group, amino group, N-methylamino group, N,N-dimethylamino group, N-ethylamino group, N,N-diethylamino group, N-propylamino group, N-isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group, N-(4-fluorophenyl)amino group, N-(pyridin-2-yl)amino group, N-(pyridin-3-yl)amino group, N-(pyridin-4-yl)amino group, N-(furan-2-yl)amino group, N-(furan-3-yl)amino group, N-(thiophen-2-yl)amino group, N-(thiophen-3-yl)amino group, pyrrolidino group, piperidino group, morpholino group, methyloxycarbonylamino group, and ethyloxycarbonylamino group. Rz' represents butyl group, isobutyl group,

2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group,  
 cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group,  
 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl  
 group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl  
 group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl  
 group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl  
 group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl  
 group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group,  
 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl  
 group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group,  
 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group,  
 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group,  
 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl  
 group, 3-methylphenylmethyl group, 4-methylphenylmethyl group,  
 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group,  
 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl  
 group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group,  
 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group,  
 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group,  
 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group,  
 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group,  
 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group,  
 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group,  
 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group,  
 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group,  
 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group,  
 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group,

2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group,  
 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,  
 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,  
 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group,  
 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group,  
 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group,  
 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group,  
 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group,  
 2-(N-phenyl-N-methylamino)ethyl group, 2-(N-ethyl-N-phenylamino)ethyl group,  
 isobutyryl group, isopropylthiocarbonyl group, isopropylsulfonyl group, valeryl  
 group, butylthiocarbonyl group, isovaleryl group, isobutylthiocarbonyl group,  
 pivaloyl group, t-butylthiocarbonyl group, cyclopropylcarbonyl group,  
 cyclopropylthiocarbonyl group, cyclopentylcarbonyl group, cyclopentylthiocarbonyl  
 group, cyclohexylcarbonyl group, cyclohexylthiocarbonyl group,  
 cyclopentylmethylcarbonyl group, cyclopentylmethylthiocarbonyl group,  
 cyclohexylmethylcarbonyl group, cyclohexylmethylthiocarbonyl group, benzoyl  
 group, thiobenzoyl group, phenylsulfonyl group, 4-methylphenylcarbonyl group,  
 4-methylphenylthiocarbonyl group, 4-methylphenylsulfonyl group,  
 4-chlorophenylcarbonyl group, 4-chlorophenylthiocarbonyl group,  
 4-fluorophenylcarbonyl group, 4-fluorophenylthiocarbonyl group,  
 isopropylloxycarbonyl group, N-isopropylcarbamoyl group, N-isopropylthiocarbamoyl  
 group, butylloxycarbonyl group, N-butylcarbamoyl group, N-butylthiocarbamoyl  
 group, isobutylloxycarbonyl group, N-isobutylcarbamoyl group,  
 N-isobutylthiocarbamoyl group, t-butylloxycarbonyl group, N-t-butylcarbamoyl  
 group, N-t-butylthiocarbamoyl group, cyclopropylloxycarbonyl group,  
 N-cyclopropylcarbamoyl group, N-cyclopropylthiocarbamoyl group,  
 cyclopentylloxycarbonyl group, N-cyclopentylcarbamoyl group,

N-cyclopentylthiocarbamoyl group, cyclohexyloxycarbonyl group, N-cyclohexylcarbamoyl group, N-cyclohexylthiocarbamoyl group, cyclopentylmethyloxycarbonyl group, cyclohexylmethyloxycarbonyl group, phenyloxycarbonyl group, N-phenylcarbamoyl group, N-phenylthiocarbamoyl group, 4-methylphenyloxycarbonyl group, N-(4-methylphenyl)carbamoyl group, N-(4-methylphenyl)thiocarbamoyl group, 4-chlorophenyloxycarbonyl group, N-(4-chlorophenyl)carbamoyl group, N-(4-chlorophenyl)thiocarbamoyl group, 4-fluorophenyloxycarbonyl group, N-(4-fluorophenyl)carbamoyl group, N-(4-fluorophenyl)thiocarbamoyl group, (pyrrolidino-1-yl)carbonyl group, (piperidino-1-yl)carbonyl group, and (morpholino-4-yl)carbonyl group. Ry' represents hydrogen atom, methyl group, ethyl group, or isobutyl group, or binds to Rz to form pyrrolidino group, piperidino group, piperazino group, morpholino group, pyrrol-1-yl group, imidazol-1-yl group, or pyrazol-1-yl group together with the nitrogen atom to which they bond. However, -D-Rx' or -N(Ry')(Rz') contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when -D-Rx' or -N(Ry')(Rz') contains amino group, the amino group may be protected with Rp<sup>2</sup>.

AR' represents any one of naphthalen-2-yl group, naphthalen-1-yl group, benzofuran-5-yl group, benzofuran-4-yl group, benzofuran-2-yl group, benzo[b]thiophen-5-yl group, benzo[b]thiophen-4-yl group, benzo[b]thiophen-2-yl group, indol-5-yl group, indol-4-yl group, indol-6-yl group, benzothiazol-6-yl group, benzothiazol-7-yl group, benzothiazol-5-yl group, benzothiazol-4-yl group, dihydro-3H-benzothiazol-6-yl group, dihydro-3H-benzothiazol-7-yl group, dihydro-3H-benzothiazol-5-yl group, dihydro-3H-benzothiazol-4-yl group, quinolin-6-yl group, quinolin-3-yl group, quinolin-5-yl group, quinolin-7-yl group, dihydro-1H-quinolin-6-yl group, dihydro-1H-quinolin-5-yl group, benzo[d]isothiazol-5-yl group, benzo[d]isothiazol-4-yl group, benzo[d]isothiazol-6-yl group, benzo[d]isothiazol-7-yl group, 1H-indazol-5-yl group, 1H-indazol-4-yl group,

1H-indazol-6-yl group, benzo[c]isothiazol-5-yl group, benzo[c]isothiazol-4-yl group,  
 benzo[c]isothiazol-6-yl group, benzo[c]isothiazol-7-yl group, 2H-indazol-5-yl group,  
 2H-indazol-4-yl group, 2H-indazol-6-yl group, imidazo[1,2-a]pyridin-6-yl group,  
 imidazo[1,2-a]pyridin-7-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1H-pyrrolo[2,3-b]pyridin-4-yl group, isoquinolin-6-yl group, isoquinolin-3-yl group,  
 isoquinolin-5-yl group, isoquinolin-7-yl group, dihydro-2H-isoquinolin-6-yl group,  
 dihydro-2H-isoquinolin-5-yl group, cinnolin-6-yl group, cinnolin-5-yl group,  
 quinazolin-6-yl group, quinazolin-7-yl group, quinazolin-5-yl group, quinoxalin-2-yl  
 group, quinoxalin-6-yl group, quinoxalin-5-yl group, 1H-benzimidazol-5-yl group,  
 1H-benzimidazol-4-yl group, benzoxazol-5-yl group, benzoxazol-6-yl group,  
 benzoxazol-4-yl group, benzoxazol-7-yl group, 1H-pyrrolo[3,2-b]pyridin-5-yl group,  
 1H-pyrrolo[3,2-b]pyridin-6-yl group, benzo[1,2,5]thiadiazol-5-yl group,  
 benzo[1,2,5]thiadiazol-4-yl group, 1H-benzotriazol-5-yl group, 1H-benzotriazol-4-yl  
 group, 1,3-dihydropyrrolo[2,3-b]pyridin-5-yl group,  
 1,3-dihydropyrrolo[2,3-b]pyridin-4-yl group, 1,3-dihydrobenzimidazol-5-yl group,  
 1,3-dihydrobenzimidazol-4-yl group, dihydro-3H-benzoxazol-6-yl group,  
 dihydro-3H-benzoxazol-7-yl group, dihydro-3H-benzoxazol-5-yl group,  
 dihydro-3H-benzoxazol-4-yl group, phthalazin-6-yl group, phthalazin-5-yl group,  
 [1,8]naphthalidin-3-yl group, [1,8]naphthalidin-4-yl group, [1,5]naphthalidin-3-yl  
 group, [1,5]naphthalidin-4-yl group, 1H-pyrrolo[3,2-c]pyridin-6-yl group,  
 1H-pyrrolo[3,2-c]pyridin-4-yl group, 1H-pyrrolo[2,3-c]pyridin-5-yl group,  
 1H-pyrrolo[2,3-c]pyridin-4-yl group, 1H-pyrazolo[4,3-b]pyridin-5-yl group,  
 1H-pyrazolo[4,3-b]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-6-yl group,  
 1H-pyrazolo[4,3-c]pyridin-4-yl group, 1H-pyrazolo[3,4-c]pyridin-5-yl group,  
 1H-pyrazolo[3,4-c]pyridin-4-yl group, 1H-pyrazolo[3,4-b]pyridin-5-yl group,  
 1H-pyrazolo[3,4-b]pyridin-4-yl group, [1,2,4]triazolo[4,3-a]pyridin-6-yl group,  
 [1,2,4]triazolo[4,3-a]pyridin-7-yl group, thieno[3,2-c]pyridin-2-yl group,

thieno[3,2-c]pyridin-3-yl group, thieno[3,2-c]pyridin-6-yl group,  
 thieno[3,2-b]pyridin-2-yl group, thieno[3,2-b]pyridin-3-yl group,  
 thieno[3,2-b]pyridin-5-yl group, thieno[3,2-b]pyridin-6-yl group,  
 1H-thieno[3,2-c]pyrazol-5-yl group, 1H-thieno[3,2-c]pyrazol-4-yl group,  
 benzo[d]isoxazol-5-yl group, benzo[d]isoxazol-4-yl group, benzo[d]isoxazol-6-yl group,  
 benzo[d]isoxazol-7-yl group, benzo[c]isoxazol-5-yl group, benzo[c]isoxazol-4-yl group,  
 benzo[c]isoxazol-6-yl group, benzo[c]isoxazol-7-yl group, indolizin-7-yl group,  
 indolizin-6-yl group, indolizine-8-yl group, 1,3-dihydroindol-5-yl group,  
 1,3-dihydroindol-4-yl group, 1,3-dihydroindol-6-yl group,  
 1H-pyrazolo[3,4-d]thiazol-5-yl group, 2H-isoindol-5-yl group, 2H-isoindol-4-yl group,  
 [1,2,4]triazolo[1,5-a]pyrimidin-6-yl group, 1H-pyrazolo[3,4-b]pyrazin-5-yl group,  
 1H-imidazo[4,5-b]pyrazin-5-yl group, 7H-purin-2-yl group, 4H-chromen-6-yl group,  
 and 4H-chromen-5-yl group (the aforementioned groups may be substituted with one  
 of Xa or two or more of identical or different Xa). The substituent Xa represents a  
 group as any one of oxo group, thioxo group, fluorine atom, chlorine atom,  
 trifluoromethyl group, methyl group, ethyl group, propyl group, 2-hydroxyethyl  
 group, carboxymethyl group, 2-carboxyethyl group, N,N-dimethylcarbamoylmethyl  
 group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, carboxymethyloxy  
 group, 2-carboxyethyloxy group, N,N-dimethylcarbamoylmethyloxy group, amino  
 group, methylamino group, dimethylamino group, 2-hydroxyethylamino group,  
 carbamoylamino group, acetylamino group, furan-2-carboxyamino group,  
 2-hydroxyacetylamino group, 2-aminoacetylamino group, methylsulfonylamino  
 group, (N,N-dimethylsulfamoyl)amino group, methanesulfonyl group, sulfamoyl  
 group, N-methylsulfamoyl group, N,N-dimethylsulfamoyl group, carboxyl group,  
 acetyl group, carbamoyl group, and N,N-dimethylcarbamoyl group. Provided that  
 when AR' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>,  
 and when substituted AR' contains amino group, the amino group may be protected

with  $Rp^2$ .

In another preferred embodiment, the compound represented by the formula (III) satisfies all of the following requirements.

$AR'$  binds to  $C^{3'}$ ,  $Rs'$  binds to any of the ring-constituting carbon atoms  $C^{4'}$ ,  $C^{5'}$ , and  $C^{6'}$ , and a ring-constituting carbon atom to which  $Rs'$  does not bind among  $C^{4'}$ ,  $C^{5'}$ , and  $C^{6'}$  may be replaced with  $V'$ .

$V'$  represents nitrogen atom, or carbon atom substituted with  $Zx'$ , and  $Zx'$  represents any one of fluorine atom, chlorine atom, bromine atom, nitro group, methyl group, hydroxyl group, methoxy group, amino group, N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, and N,N-dimethylsulfamoylamino group, provided that when  $Zx'$  contains hydroxyl group, the hydroxyl group may be protected with  $Rp^1$ , and when the substituted  $Zx'$  contains amino group, the amino group may be protected with  $Rp^2$ .

$Rs'$  represents  $-D-Rx'$  or  $-N(Ry')(Rz')$ .  $D$  represents oxygen atom or sulfur atom.  $Rx'$  represents butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-cyclopentylethyl group, or 2-cyclohexylethyl group, or represents  $Rb$  or  $Rc$ .  $Q$  in  $Rb$  represents a group as any one of phenyl group, thienyl group, furyl group, pyridyl group, oxazolyl group, naphthyl group, tetrahydronaphthyl group, indanyl group, indolyl group, and dihydrobenzodioxyl group.  $A^2$  represents a single bond, oxygen atom, sulfur atom,  $-N(methyl)-$ , or  $-N(ethyl)-$  (provided that when  $A^2$  represents oxygen atom, sulfur atom,  $-N(methyl)-$ , or  $-N(ethyl)-$ ,  $A^1$  represents ethylene).  $R^2$  and  $R^3$  independently represent hydrogen atom, methyl group, fluorine atom, chlorine atom, trifluoromethyl group, methoxy group, dimethylamino group, acetylamino group, or methylsulfonylamino group



(provided that when Q represents phenyl group, A<sup>1</sup> represents a single bond, or unsubstituted methylene, and A<sup>2</sup> represents a single bond, one of R<sup>2</sup> and R<sup>3</sup> represents a substituent other than hydrogen atom). Symbol p in R<sub>c</sub> represents an integer of 2 or 3, and A<sup>4</sup> represents a single bond or methylene. A<sup>5</sup> represents -C(O)-, -C(S)-, or -S(O)<sub>2</sub>-. R<sub>d</sub> represents hydrogen atom, or a group as any one of methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group, cyclopropylmethyl group, cyclopentyl group, cyclopentylmethyl group, cyclohexyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, and pyridin-4-yl group. R<sub>e</sub> represents any one of methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, phenylmethyl group, 4-chlorophenylmethyl group, 4-fluorophenylmethyl group, pyridin-2-yl group, pyridin-3-yl group, pyridin-4-yl group, furan-2-yl group, furan-3-yl group, thiophen-2-yl group, thiophen-3-yl group, methoxy group, ethoxy group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group, phenyloxy group, 4-methylphenyloxy group, 4-chlorophenyloxy group, 4-fluorophenyloxy group, thiomethoxy group, amino group, N-methylamino group, N,N-dimethylamino group, N-ethylamino group, N,N-diethylamino group, N-propylamino group, N-isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group,

N-(4-fluorophenyl)amino group, N-(pyridin-2-yl)amino group, N-(pyridin-3-yl)amino group, N-(pyridin-4-yl)amino group, N-(furan-2-yl)amino group, N-(furan-3-yl)amino group, N-(thiophen-2-yl)amino group, N-(thiophen-3-yl)amino group, pyrrolidino group, piperidino group, morpholino group, methyloxycarbonylamino group, and ethyloxycarbonylamino group. Rz' represents any one of butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group,

3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group,  
 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group,  
 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group,  
 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group,  
 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group,  
 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group,  
 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,  
 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,  
 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group,  
 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group,  
 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group,  
 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group,  
 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group,  
 2-(N-phenyl-N-methylamino)ethyl group, 2-(N-ethyl-N-phenylamino)ethyl group,  
 isobutyryl group, isopropylthiocarbonyl group, isopropylsulfonyl group, valeryl  
 group, butylthiocarbonyl group, isovaleryl group, isobutylthiocarbonyl group,  
 pivaloyl group, t-butylthiocarbonyl group, cyclopropylcarbonyl group,  
 cyclopropylthiocarbonyl group, cyclopentylcarbonyl group, cyclopentylthiocarbonyl  
 group, cyclohexylcarbonyl group, cyclohexylthiocarbonyl group,  
 cyclopentylmethylcarbonyl group, cyclopentylmethylthiocarbonyl group,  
 cyclohexylmethylcarbonyl group, cyclohexylmethylthiocarbonyl group, benzoyl  
 group, thiobenzoyl group, phenylsulfonyl group, 4-methylphenylcarbonyl group,  
 4-methylphenylthiocarbonyl group, 4-methylphenylsulfonyl group,  
 4-chlorophenylcarbonyl group, 4-chlorophenylthiocarbonyl group,  
 4-fluorophenylcarbonyl group, 4-fluorophenylthiocarbonyl group,  
 isopropylloxycarbonyl group, N-isopropylcarbamoyl group, N-isopropylthiocarbamoyl  
 group, butylloxycarbonyl group, N-butylcarbamoyl group, N-butylthiocarbamoyl

group, isobutyloxycarbonyl group, N-isobutylcarbamoyl group, N-isobutylthiocarbamoyl group, t-butyloxycarbonyl group, N-t-butylcarbamoyl group, N-t-butylthiocarbamoyl group, cyclopropyloxycarbonyl group, N-cyclopropylcarbamoyl group, N-cyclopropylthiocarbamoyl group, cyclopentyloxycarbonyl group, N-cyclopentylcarbamoyl group, N-cyclopentylthiocarbamoyl group, cyclohexyloxycarbonyl group, N-cyclohexylcarbamoyl group, N-cyclohexylthiocarbamoyl group, cyclopentylmethyloxycarbonyl group, cyclohexylmethyloxycarbonyl group, phenyloxycarbonyl group, N-phenylcarbamoyl group, N-phenylthiocarbamoyl group, 4-methylphenyloxycarbonyl group, N-(4-methylphenyl)carbamoyl group, N-(4-methylphenyl)thiocarbamoyl group, 4-chlorophenyloxycarbonyl group, N-(4-chlorophenyl)carbamoyl group, N-(4-chlorophenyl)thiocarbamoyl group, 4-fluorophenyloxycarbonyl group, N-(4-fluorophenyl)carbamoyl group, N-(4-fluorophenyl)thiocarbamoyl group, (pyrrolidino-1-yl)carbonyl group, (piperidino-1-yl)carbonyl group, and (morpholino-4-yl)carbonyl group.  $R_y'$  represents hydrogen atom, methyl group, ethyl group, or isobutyl group, or binds to  $R_z'$  to form pyrrolidino group, piperidino group, piperazino group, morpholino group, pyrrol-1-yl group, imidazol-1-yl group, or pyrazol-1-yl group together with nitrogen atom. Provided that when  $-D-R_x'$  or  $-N(R_y')(R_z')$  contains hydroxyl group, the hydroxyl group may be protected with  $R_p^1$ , and when  $-D-R_x'$  or  $-N(R_y')(R_z')$  contains amino group, the amino group may be protected with  $R_p^2$ .

$AR'$  represents any one of naphthalen-2-yl group, naphthalen-1-yl group, benzofuran-5-yl group, benzofuran-4-yl group, benzofuran-2-yl group, benzo[b]thiophen-5-yl group, benzo[b]thiophen-4-yl group, benzo[b]thiophen-2-yl group, indol-5-yl group, indol-4-yl group, indol-6-yl group, benzothiazol-6-yl group, benzothiazol-7-yl group, benzothiazol-5-yl group, benzothiazol-4-yl group, dihydro-3H-benzothiazol-6-yl group, dihydro-3H-benzothiazol-7-yl group,

dihydro-3H-benzothiazol-5-yl group, dihydro-3H-benzothiazol-4-yl group,  
 quinolin-6-yl group, quinolin-3-yl group, quinolin-5-yl group, quinolin-7-yl group,  
 dihydro-1H-quinolin-6-yl group, dihydro-1H-quinolin-5-yl group,  
 benzo[d]isothiazol-5-yl group, benzo[d]isothiazol-4-yl group, benzo[d]isothiazol-6-yl  
 group, benzo[d]isothiazol-7-yl group, 1H-indazol-5-yl group, 1H-indazol-4-yl group,  
 1H-indazol-6-yl group, benzo[c]isothiazol-5-yl group, benzo[c]isothiazol-4-yl group,  
 benzo[c]isothiazol-6-yl group, benzo[c]isothiazol-7-yl group, 2H-indazol-5-yl group,  
 2H-indazol-4-yl group, 2H-indazol-6-yl group, imidazo[1,2-a]pyridin-6-yl group,  
 imidazo[1,2-a]pyridin-7-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1H-pyrrolo[2,3-b]pyridin-4-yl group, isoquinolin-6-yl group, isoquinolin-3-yl group,  
 isoquinolin-5-yl group, isoquinolin-7-yl group, dihydro-2H-isoquinolin-6-yl group,  
 dihydro-2H-isoquinolin-5-yl group, cinnolin-6-yl group, cinnolin-5-yl group,  
 quinazolin-6-yl group, quinazolin-7-yl group, quinazolin-5-yl group, quinoxalin-2-yl  
 group, quinoxalin-6-yl group, quinoxalin-5-yl group, 1H-benzimidazol-5-yl group,  
 1H-benzimidazol-4-yl group, benzoxazol-5-yl group, benzoxazol-6-yl group,  
 benzoxazol-4-yl group, benzoxazol-7-yl group, 1H-pyrrolo[3,2-b]pyridin-5-yl group,  
 1H-pyrrolo[3,2-b]pyridin-6-yl group, benzo[1,2,5]thiadiazol-5-yl group,  
 benzo[1,2,5]thiadiazol-4-yl group, 1H-benzotriazol-5-yl group, 1H-benzotriazol-4-yl  
 group, 1,3-dihydropyrrolo[2,3-b]pyridin-5-yl group,  
 1,3-dihydropyrrolo[2,3-b]pyridin-4-yl group, 1,3-dihydrobenzimidazol-5-yl group,  
 1,3-dihydrobenzimidazol-4-yl group, dihydro-3H-benzoxazol-6-yl group,  
 dihydro-3H-benzoxazol-7-yl group, dihydro-3H-benzoxazol-5-yl group,  
 dihydro-3H-benzoxazol-4-yl group, phthalazin-6-yl group, phthalazin-5-yl group,  
 [1,8]naphthalidin-3-yl group, [1,8]naphthalidin-4-yl group, [1,5]naphthalidin-3-yl  
 group, [1,5]naphthalidin-4-yl group, 1H-pyrrolo[3,2-c]pyridin-6-yl group,  
 1H-pyrrolo[3,2-c]pyridin-4-yl group, 1H-pyrrolo[2,3-c]pyridin-5-yl group,  
 1H-pyrrolo[2,3-c]pyridin-4-yl group, 1H-pyrazolo[4,3-b]pyridin-5-yl group,

1H-pyrazolo[4,3-b]pyridin-6-yl group, 1H-pyrazolo[4,3-c]pyridin-6-yl group,  
 1H-pyrazolo[4,3-c]pyridin-4-yl group, 1H-pyrazolo[3,4-c]pyridin-5-yl group,  
 1H-pyrazolo[3,4-c]pyridin-4-yl group, 1H-pyrazolo[3,4-b]pyridin-5-yl group,  
 1H-pyrazolo[3,4-b]pyridin-4-yl group, [1,2,4]triazolo[4,3-a]pyridin-6-yl group,  
 [1,2,4]triazolo[4,3-a]pyridin-7-yl group, thieno[3,2-c]pyridin-2-yl group,  
 thieno[3,2-c]pyridin-3-yl group, thieno[3,2-c]pyridin-6-yl group,  
 thieno[3,2-b]pyridin-2-yl group, thieno[3,2-b]pyridin-3-yl group,  
 thieno[3,2-b]pyridin-5-yl group, thieno[3,2-b]pyridin-6-yl group,  
 1H-thieno[3,2-c]pyrazol-5-yl group, 1H-thieno[3,2-c]pyrazol-4-yl group,  
 benzo[d]isoxazol-5-yl group, benzo[d]isoxazol-4-yl group, benzo[d]isoxazol-6-yl group,  
 benzo[d]isoxazol-7-yl group, benzo[c]isoxazol-5-yl group, benzo[c]isoxazol-4-yl group,  
 benzo[c]isoxazol-6-yl group, benzo[c]isoxazol-7-yl group, indolizin-7-yl group,  
 indolizin-6-yl group, indolizine-8-yl group, 1,3-dihydroindol-5-yl group,  
 1,3-dihydroindol-4-yl group, 1,3-dihydroindol-6-yl group,  
 1H-pyrazolo[3,4-d]thiazol-5-yl group, 2H-isoindol-5-yl group, 2H-isoindol-4-yl group,  
 [1,2,4]triazolo[1,5-a]pyrimidin-6-yl group, 1H-pyrazolo[3,4-b]pyrazin-5-yl group,  
 1H-imidazo[4,5-b]pyrazin-5-yl group, 7H-purin-2-yl group, 4H-chromen-6-yl group,  
 and 4H-chromen-5-yl group (the aforementioned groups may be substituted with one  
 of Xa or two or more of identical or different Xa). The substituent Xa represents a  
 group as any one of oxo group, thioxo group, fluorine atom, chlorine atom,  
 trifluoromethyl group, methyl group, ethyl group, propyl group, 2-hydroxyethyl  
 group, carboxymethyl group, 2-carboxyethyl group, N,N-dimethylcarbamoylmethyl  
 group, hydroxyl group, methoxy group, 2-hydroxyethyloxy group, carboxymethyloxy  
 group, 2-carboxyethyloxy group, N,N-dimethylcarbamoylmethyloxy group, amino  
 group, methylamino group, dimethylamino group, 2-hydroxyethylamino group,  
 carbamoylamino group, acetylamino group, furan-2-carboxyamino group,  
 2-hydroxyacetylamino group, 2-aminoacetylamino group, methylsulfonylamino

group, (N,N-dimethylsulfamoyl)amino group, methanesulfonyl group, sulfamoyl group, N-methylsulfamoyl group, N,N-dimethylsulfamoyl group, carboxyl group, acetyl group, carbamoyl group, and N,N-dimethylcarbamoyl group. Provided that when AR' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when substituted AR' contains amino group, the amino group may be protected with Rp<sup>2</sup>.

In a particularly preferred embodiment, the compound represented by the formula (III) satisfies all of the following requirements.

C<sup>2'</sup> represents carbon atom to which AR' binds, C<sup>3'</sup> represents carbon atom to which Rs' binds, C<sup>4'</sup> may be replaced with V', and C<sup>5'</sup> and C<sup>6'</sup> represent an unsubstituted ring-constituting carbon atom.

V' represents nitrogen atom, or carbon atom substituted with Zx', and Zx' represents any one of fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, and N,N-dimethylamino group, provided that when Zx' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when the substituted Zx' contains amino group, the amino group may be protected with Rp<sup>2</sup>.

Rs' represents -O-Rx'. Rx' represents any one of butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl

group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group,  
 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group,  
 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group,  
 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl  
 group, 3-methylphenylmethyl group, 4-methylphenylmethyl group,  
 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group,  
 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl  
 group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group,  
 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group,  
 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group,  
 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group,  
 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group,  
 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group,  
 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group,  
 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group,  
 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group,  
 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group,  
 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group,  
 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group,  
 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,  
 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,  
 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group,  
 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group,  
 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group,  
 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group,  
 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group,  
 2-(N-phenyl-N-methylamino)ethyl group, and 2-(N-ethyl-N-phenylamino)ethyl



group.

AR' represents any one of naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethoxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group, 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl

group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, and benzoxazol-5-yl group. Provided that when AR' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when substituted AR' contains amino group, the amino group may be protected with Rp<sup>2</sup>.

In another particularly preferred embodiment, the compound represented by the formula (III) satisfies all of the following requirements.

C<sup>2'</sup> represents carbon atom to which AR' binds, C<sup>4'</sup> represents carbon atom to which Rs' binds, C<sup>5'</sup> may be replaced with V', and C<sup>3'</sup> and C<sup>6'</sup> represent an unsubstituted ring-constituting carbon atom.

V' represents nitrogen atom, or carbon atom substituted with Zx', and Zx' represents any one of fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, and N,N-dimethylamino group. Provided that when Zx' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when the substituted Zx' contains amino group, the amino group may be protected with Rp<sup>2</sup>.

Rs' represents -O-Rx'. Rx' represents any one of butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl

group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl  
 group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl  
 group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl  
 group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl  
 group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group,  
 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl  
 group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group,  
 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group,  
 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group,  
 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl  
 group, 3-methylphenylmethyl group, 4-methylphenylmethyl group,  
 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group,  
 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl  
 group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group,  
 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group,  
 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group,  
 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group,  
 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group,  
 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group,  
 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group,  
 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group,  
 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group,  
 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group,  
 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group,  
 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group,  
 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,  
 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,

2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, and 2-(N-ethyl-N-phenylamino)ethyl group.

AR' represents any one of naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group,

2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group,  
 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group,  
 quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl  
 group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group,  
 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group,  
 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group,  
 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group,  
 imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl  
 group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group,  
 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, and benzoxazol-5-yl  
 group. Provided that when AR' contains hydroxyl group, the hydroxyl group may be  
 protected with Rp<sup>1</sup>, and when substituted AR' contains amino group, the amino  
 group may be protected with Rp<sup>2</sup>.

In another particularly preferred embodiment, the compound represented by  
 the formula (III) satisfies all of the following requirements.

C<sup>3'</sup> represents carbon atom to which AR' binds, C<sup>5'</sup> represents carbon atom to  
 which Rs' binds, and C<sup>2'</sup>, C<sup>4'</sup> and C<sup>6'</sup> represent an unsubstituted ring-constituting  
 carbon atom.

Rs' represents -O-Rx'. Rx' represents any one of butyl group, isobutyl group,  
 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group,  
 cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group,  
 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl

group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl  
 group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl  
 group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl  
 group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl  
 group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group,  
 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl  
 group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group,  
 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group,  
 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group,  
 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl  
 group, 3-methylphenylmethyl group, 4-methylphenylmethyl group,  
 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group,  
 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl  
 group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group,  
 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group,  
 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group,  
 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group,  
 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group,  
 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group,  
 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group,  
 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group,  
 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group,  
 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group,  
 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group,  
 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group,  
 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,  
 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,

2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group, 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group, 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group, 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group, 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group, 2-(N-phenyl-N-methylamino)ethyl group, and 2-(N-ethyl-N-phenylamino)ethyl group.

AR' represents any one of naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group,

2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, and benzoxazol-5-yl group. Provided that when AR' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when substituted AR' contains amino group, the amino group may be protected with Rp<sup>2</sup>.

In another particularly preferred embodiment, the compound represented by the formula (III) satisfies all of the following requirements.

C<sup>3'</sup> represents carbon atom to which AR' binds, C<sup>4'</sup> represents carbon atom to which Rs' binds, C<sup>5'</sup> represents nitrogen atom, and C<sup>2'</sup> and C<sup>6'</sup> represent an unsubstituted ring-constituting carbon atom.

Rs' represents -O-Rx'. Rx' represents any one of butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl



group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl  
 group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl  
 group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl  
 group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl  
 group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group,  
 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl  
 group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group,  
 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group,  
 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group,  
 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl  
 group, 3-methylphenylmethyl group, 4-methylphenylmethyl group,  
 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group,  
 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl  
 group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group,  
 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group,  
 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group,  
 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group,  
 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group,  
 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group,  
 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group,  
 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group,  
 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group,  
 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group,  
 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group,  
 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group,  
 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,  
 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,

2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group,  
 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group,  
 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group,  
 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group,  
 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group,  
 2-(N-phenyl-N-methylamino)ethyl group, and 2-(N-ethyl-N-phenylamino)ethyl  
 group.

AR' represents any one of naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl  
 group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl  
 group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group,  
 6-(N,N-dimethylamino)naphthalen-2-yl group,  
 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group,  
 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group,  
 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group,  
 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group,  
 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group,  
 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group,  
 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group,  
 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group,  
 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group,  
 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group,  
 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group,  
 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group,  
 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group,  
 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group,

2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group,  
 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group,  
 quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl  
 group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group,  
 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group,  
 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group,  
 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group,  
 imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl  
 group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group,  
 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, and benzoxazol-5-yl  
 group. Provided that when AR' contains hydroxyl group, the hydroxyl group may be  
 protected with Rp<sup>1</sup>, and when substituted AR' contains amino group, the amino  
 group may be protected with Rp<sup>2</sup>.

In another particularly preferred embodiment, the compound represented by  
 the formula (III) satisfies all of the following requirements.

C<sup>3'</sup> represents carbon atom to which AR' binds, C<sup>4'</sup> represents carbon atom to  
 which Rs' binds, C<sup>6'</sup> represents carbon atom substituted with Zx', and C<sup>2'</sup> and C<sup>5'</sup>  
 represent an unsubstituted ring-constituting carbon atom.

Zx' represents any one of fluorine atom, methyl group, hydroxyl group,  
 amino group, N-methylamino group, and N,N-dimethylamino group, provided that  
 when Zx' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>,  
 and when the substituted Zx' contains amino group, the amino group may be

protected with  $R_p^2$ .

$R_s'$  represents  $-O-R_x'$ .  $R_x'$  represents any one of butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group, 3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group, 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group, 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group, 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group, 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group, 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group,

2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group,  
 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group,  
 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group,  
 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,  
 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,  
 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group,  
 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group,  
 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group,  
 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group,  
 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group,  
 2-(N-phenyl-N-methylamino)ethyl group, and 2-(N-ethyl-N-phenylamino)ethyl group.

AR' represents any one of naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group,

2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group, 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group, 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group, 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-2,3-dihydrobenzothiazol-6-yl group, 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group, quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group, 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group, 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group, 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group, imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group, 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, and benzoxazol-5-yl group. Provided that when AR' contains hydroxyl group, the hydroxyl group may be protected with Rp<sup>1</sup>, and when substituted AR' contains amino group, the amino group may be protected with Rp<sup>2</sup>.

In another particularly preferred embodiment, the compound represented by the formula (III) satisfies all of the following requirements..

C<sup>3'</sup> represents carbon atom to which AR' binds, C<sup>4'</sup> represents carbon atom to which Rs' binds, C<sup>5'</sup> represents carbon atom substituted with Zx', and C<sup>2'</sup> and C<sup>6'</sup>

represent an unsubstituted ring-constituting carbon atom.

Zx' represents any one of N-methylamino group, N-ethylamino group, N-propylamino group, N-isopropylamino group, N,N-dimethylamino group, N,N-diethylamino group, formylamino group, acetylamino group, carbamoylamino group, mesylamino group, and N,N-dimethylsulfamoylamino group. Provided that when the substituted Zx' contains amino group (NH), the amino group may be protected with Rp<sup>2</sup>.

Rs' represents -O-Rx'. Rx' represents any one of butyl group, isobutyl group, 2-ethylbutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclopentylmethyl group, cyclohexylmethyl group, 2-methylphenyl group, 4-methylphenyl group, 2-fluorophenyl group, 3-fluorophenyl group, 4-fluorophenyl group, 2-chlorophenyl group, 3-chlorophenyl group, 4-chlorophenyl group, indan-2-yl group, 4-methylindan-2-yl group, 5-methylindan-2-yl group, 4,7-dimethylindan-2-yl group, 5,6-dimethylindan-2-yl group, 4-fluoroindan-2-yl group, 5-fluoroindan-2-yl group, 4,7-difluoroindan-2-yl group, 5,6-difluoroindan-2-yl group, 4-chloroindan-2-yl group, 5-chloroindan-2-yl group, 4,7-dichloroindan-2-yl group, 5,6-dichloroindan-2-yl group, 4-methoxyindan-2-yl group, 5-methoxyindan-2-yl group, 4,7-dimethoxyindan-2-yl group, 5,6-dimethoxyindan-2-yl group, 1-phenylethyl group, 1-(2-fluorophenyl)ethyl group, 1-(3-fluorophenyl)ethyl group, 1-(4-fluorophenyl)ethyl group, 1-(2-chlorophenyl)ethyl group, 1-(3-chlorophenyl)ethyl group, 1-(4-chlorophenyl)ethyl group, 2-methylphenylmethyl group, 3-methylphenylmethyl group, 4-methylphenylmethyl group, 2,3-dimethylphenylmethyl group, 3,5-dimethylphenylmethyl group, 2-fluorophenylmethyl group, 3-fluorophenylmethyl group, 4-fluorophenylmethyl group, 2-chlorophenylmethyl group, 3-chlorophenylmethyl group, 4-chlorophenylmethyl group, 2,3-difluorophenylmethyl group, 2,4-difluorophenylmethyl group, 2,5-difluorophenylmethyl group,

3,4-difluorophenylmethyl group, 2,3-dichlorophenylmethyl group,  
 2,4-dichlorophenylmethyl group, 2,5-dichlorophenylmethyl group,  
 2,6-dichlorophenylmethyl group, 3,4-dichlorophenylmethyl group,  
 3,5-dichlorophenylmethyl group, 3,6-dichlorophenylmethyl group,  
 2-(trifluoromethyl)phenylmethyl group, 3-(trifluoromethyl)phenylmethyl group,  
 4-(trifluoromethyl)phenylmethyl group, 2-(2-methylphenyl)ethyl group,  
 2-(3-methylphenyl)ethyl group, 2-(4-methylphenyl)ethyl group,  
 2-(2-methoxyphenyl)ethyl group, 2-(3-methoxyphenyl)ethyl group,  
 2-(4-methoxyphenyl)ethyl group, 2-(2-fluorophenyl)ethyl group,  
 2-(3-fluorophenyl)ethyl group, 2-(4-fluorophenyl)ethyl group,  
 2-(2-chlorophenyl)ethyl group, 2-(3-chlorophenyl)ethyl group,  
 2-(4-chlorophenyl)ethyl group, 2-[2-(trifluoromethyl)phenyl]ethyl group,  
 2-[3-(trifluoromethyl)phenyl]ethyl group, 2-[4-(trifluoromethyl)phenyl]ethyl group,  
 2-[4-(N,N-dimethylamino)phenyl]ethyl group, 2-phenyloxyethyl group,  
 2-(2-chlorophenyloxy)ethyl group, 2-(3-chlorophenyloxy)ethyl group,  
 2-(4-chlorophenyloxy)ethyl group, 2-(phenylthio)ethyl group,  
 2-(N-phenyl-N-methylamino)ethyl group, and 2-(N-ethyl-N-phenylamino)ethyl group.

AR' represents any one of naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group,



2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group,  
 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group,  
 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group,  
 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group,  
 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group,  
 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group,  
 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group,  
 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group,  
 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group,  
 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group,  
 2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group,  
 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group,  
 quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl  
 group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group,  
 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group,  
 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group,  
 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group,  
 imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl  
 group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group,  
 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, and benzoxazol-5-yl  
 group. Provided that when AR' contains hydroxyl group, the hydroxyl group may be

protected with  $Rp^1$ , and when substituted  $AR'$  contains amino group, the amino group may be protected with  $Rp^2$ .

In another particularly preferred embodiment, the compound represented by the formula (III) satisfies all of the following requirements.

$C^{3'}$  represents carbon atom to which  $AR'$  binds,  $C^{4'}$  represents carbon atom to which  $Rs'$  binds,  $C^{5'}$  represents carbon atom substituted with  $Zx'$ , or an unsubstituted ring-constituting carbon atom, and  $C^{2'}$  and  $C^{6'}$  represent an unsubstituted ring-constituting carbon atom.

$Zx'$  represents any one of fluorine atom, methyl group, hydroxyl group, amino group, N-methylamino group, and N,N-dimethylamino group, provided that when  $Zx'$  contains hydroxyl group, the hydroxyl group may be protected with  $Rp^1$ , and when the substituted  $Zx'$  contains amino group, the amino group may be protected with  $Rp^2$ .

$Rs'$  represents  $-O-Rx'$ .  $Rx'$  have the same meaning as that of  $Rc$ , provided that when  $Rc$  contains hydroxyl group, the hydroxyl group may be protected with  $Rp^1$ .  $p$  in  $Rc$  represents an integer of 2, and  $A^4$  represents a single bond or methylene.  $A^5$  represents  $-C(O)-$ ,  $-C(S)-$ , or  $-S(O)_2-$ .  $Rd$  represents a group as any one of methyl group, ethyl group, propyl group, isopropyl group, butyl group, isobutyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, benzyl group, 4-chlorophenylmethyl group, and 4-fluorophenylmethyl group.  $Re$  represents a group as any one of isopropyl group, butyl group, isobutyl group, t-butyl group, cyclopropyl group, cyclopentyl group, cyclohexyl group, cyclopentylmethyl group, cyclohexylmethyl group, phenyl group, 4-methylphenyl group, 4-chlorophenyl group, 4-fluorophenyl group, propyloxy group, isopropyloxy group, butyloxy group, isobutyloxy group, t-butyloxy group, cyclopropyloxy group, cyclopentyloxy group, cyclohexyloxy group, cyclopentylmethyloxy group, cyclohexylmethyloxy group,

phenyloxy group, 4-methylphenyloxy group, 4-chlorophenyloxy group, 4-fluorophenyloxy group, N-propylamino group, N-isopropylamino group, N-butylamino group, N-isobutylamino group, N-t-butylamino group, N-cyclopropylamino group, N-cyclopentylamino group, N-cyclohexylamino group, N-phenylamino group, N-(4-methylphenyl)amino group, N-(4-chlorophenyl)amino group, N-(4-fluorophenyl)amino group, pyrrolidino group, piperidino group, and morpholino group.

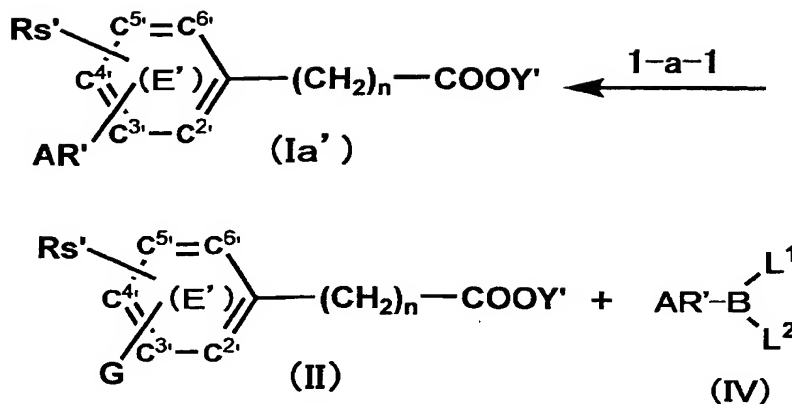
AR' represents any one of naphthalen-2-yl group, 6-hydroxynaphthalen-2-yl group, 6-methoxynaphthalen-2-yl group, 6-(2-hydroxyethyloxy)naphthalen-2-yl group, 6-aminonaphthalen-2-yl group, 6-(N-methylamino)naphthalen-2-yl group, 6-(N,N-dimethylamino)naphthalen-2-yl group, 6-(2-hydroxyethylamino)naphthalen-2-yl group, benzo[b]furan-5-yl group, 2-methylbenzo[b]furan-5-yl group, 3-methylbenzo[b]furan-5-yl group, 2,3-dimethylbenzo[b]furan-5-yl group, benzo[b]thiophen-5-yl group, 2-methylbenzo[b]thiophen-5-yl group, 3-methylbenzo[b]thiophen-5-yl group, 2,3-dimethylbenzo[b]thiophen-5-yl group, 1H-indol-5-yl group, 2-methyl-1H-indol-5-yl group, 3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1H-indol-5-yl group, 1-methyl-1H-indol-5-yl group, 1,2-dimethyl-1H-indol-5-yl group, 1,3-dimethyl-1H-indol-5-yl group, 1,2,3-trimethyl-1H-indol-5-yl group, 1-ethyl-1H-indol-5-yl group, 1-ethyl-2-methyl-1H-indol-5-yl group, 1-ethyl-3-methyl-1H-indol-5-yl group, 1-ethyl-2,3-dimethyl-1H-indol-5-yl group, 1-propyl-1H-indol-5-yl group, 2-methyl-1-propyl-1H-indol-5-yl group, 3-methyl-1-propyl-1H-indol-5-yl group, 2,3-dimethyl-1-propyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-1H-indol-5-yl group, 1-(2-hydroxyethyl)-2-methyl-1H-indol-5-yl group, 1-(2-hydroxyethyl)-3-methyl-1H-indol-5-yl group, 2,3-dimethyl-1-(2-hydroxyethyl)-1H-indol-5-yl group, benzothiazol-6-yl group,

2-methylbenzothiazol-6-yl group, 2-methoxybenzothiazol-6-yl group,  
 2-aminobenzothiazol-6-yl group, 2-oxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-oxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-2,3-dihydrobenzothiazol-6-yl group,  
 2-thioxo-3-methyl-2,3-dihydrobenzothiazol-6-yl group, quinolin-3-yl group,  
 quinolin-6-yl group, 2-oxo-1,2-dihydroquinolin-6-yl group, benzo[d]isothiazol-5-yl  
 group, 1H-indazol-5-yl group, 1-methyl-1H-indazol-5-yl group,  
 1-ethyl-1H-indazol-5-yl group, 1-propyl-1H-indazol-5-yl group,  
 1-(2-hydroxyethyl)-1H-indazol-5-yl group, 3-hydroxy-1H-indazol-5-yl group,  
 3-hydroxy-1-methyl-1H-indazol-5-yl group, 1-ethyl-3-hydroxy-1H-indazol-5-yl group,  
 imidazo[1,2-a]pyridin-6-yl group, 1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-methyl-1H-pyrrolo[2,3-b]pyridin-5-yl group, 1-ethyl-1H-pyrrolo[2,3-b]pyridin-5-yl  
 group, 1-propyl-1H-pyrrolo[2,3-b]pyridin-5-yl group,  
 1-(2-hydroxyethyl)-1H-pyrrolo[2,3-b]pyridin-5-yl group, isoquinolin-6-yl group,  
 1-oxo-1,2-dihydroisoquinolin-6-yl group, cinnolin-6-yl group, and benzoxazol-5-yl  
 group. Provided that when AR' contains hydroxyl group, the hydroxyl group may be  
 protected with Rp<sup>1</sup>, and when substituted AR' contains amino group, the amino  
 group may be protected with Rp<sup>2</sup>.

Compound (I) of the present invention can be produced by, for example,  
 employing the reactions according to the following various methods.

[Preparation Method 1] (Step a-1)

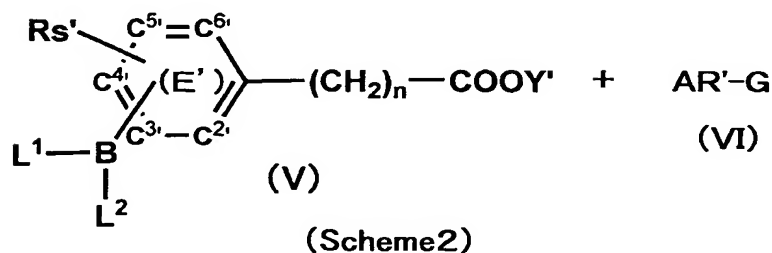
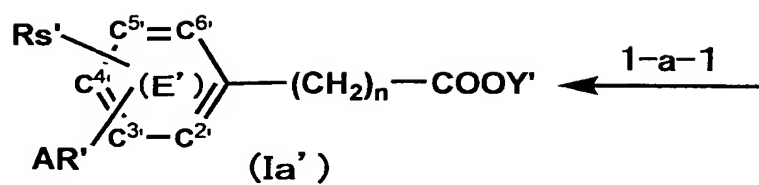
As shown in the following scheme 1:



(Scheme 1)

a compound of the present invention represented by the formula (Ia') wherein Y represents a lower alkyl group having 1 to 4 carbon atoms, and Rs, AR, and V on or in the aromatic ring (E) may be protected [hereinafter simply referred to as "Compound (Ia')"], which falls within the scope of Compound (I) of the present invention, can be prepared by reacting a compound represented by the formula (II) [simply referred to as "Compound (II)" hereinafter] with a boronic acid derivative represented by the formula (IV) [hereinafter simply referred to as "Compound (IV)"]. n, C<sup>2'</sup> to C<sup>6'</sup>, Rs', AR', Y' and G in the formulas have the same meanings as defined above. In the formula of Compound (IV), L<sup>1</sup> and L<sup>2</sup> independently represent hydroxyl group, an alkoxyl group having 1 to 8 carbon atoms (e.g., methoxy group, ethoxy group, propoxy group, isopropoxy group, cyclohexyloxy group), or a substituted or unsubstituted phenyloxy group, or L<sup>1</sup> and L<sup>2</sup> bind to each other to represent a 5- or 6-membered cyclic ester of an arylboric acid (e.g., 9-borabicyclo[3,3,1]nonane, 1,3,2-dioxaborolane, 4,4,5,5-tetramethyl-1,3,2-dioxaborolane), which forms a ring containing boron atom [this ring may be saturated or unsaturated, may be a ring containing a heteroatom other than boron (e.g., oxygen atom), and may be further substituted].

Further, as shown in the following scheme 2:



an example of the method for preparing Compound (Ia') includes a method of reacting a combination of a compound represented by the formula (V) [hereinafter simply referred to as "Compound (V)"] and a compound represented by the formula (VI) [hereinafter simply referred to as "Compound (VI)"].

Examples include a method of preparing Compound (Ia') by performing the Suzuki reaction described in, for example, Jikken Kagaku Koza, 4th Edition (edited by Chemical Society of Japan, published by Maruzen Co., Ltd.), vol. 25, p.403 with a combination mentioned either in the scheme 1 or scheme 2 or the both. A specific example includes a reaction of Compound (II) [or Compound (V)] with Compound (IV) [or Compound (VI)] in a solvent in the presence of a commercially available palladium catalyst or a catalyst prepared from a palladium complex and a ligand, and a base.

As the palladium catalyst, a commercially available catalyst such as tetrakis(triphenylphosphine)palladium, tetrakis(methyldiphenylphosphine)palladium, dichlorobis(triphenylphosphine)palladium, dichlorobis(tri-*o*-tolylphosphine)palladium, dichlorobis(tricyclohexylphosphine)palladium, dichlorobis(triethylphosphine)palladium, palladium acetate, palladium chloride,

bis(acetonitrile)palladium chloride, tris(dibenzylideneacetone)dipalladium and bis(diphenylphosphinoferrocene)palladium chloride may be purchased and added to the reaction system, per se, or a catalyst may be added which is separately prepared from palladium acetate, tris(dibenzylideneacetone)dipalladium or the like and arbitrary ligands and isolated. Further, a catalyst considered to actually participate in the reaction may also be prepared by mixing palladium acetate, tris(dibenzylideneacetone)dipalladium or the like and arbitrary ligands in the reaction system. The valence of palladium may be 0 or may be +2. Examples of the ligand include phosphine ligands such as trifurylphosphine, tri(o-tolyl)phosphine, tri(cyclohexyl)phosphine, tri(t-butyl)phosphine, dicyclohexylphenylphosphine, 1,1'-bis(di-t-butylphosphino)ferrocene, 2-dicyclohexylphosphino-2'-dimethylamino-1,1'-biphenyl and 2-(di-t-butylphosphino)biphenyl and phosphine mimic ligands such as imidazol-2-ylidenecarbenes. Chemical equivalents of the palladium catalyst may be one equivalent or a catalytic amount, and the amount may preferably be 0.01 to 20.0 mol %, and most preferably be 0.10 to 10.0 mol %.

Examples of the base include sodium carbonate, potassium carbonate, cesium carbonate, cesium fluoride, potassium fluoride, potassium phosphate, potassium acetate, triethylamine, potassium hydroxide, sodium hydroxide, sodium methoxide, lithium methoxide and the like. The reaction temperature is, for example, preferably 20°C to 150°C, and particularly preferable examples include 20°C to 120°C.

The reaction system may be either a two-phase system of water and an organic solvent, or a homogeneous system of a water-containing organic solvent or an organic solvent. As for the organic solvent, examples include uses of hydrocarbon-type solvents such as toluene, xylene and hexane, halogen-type solvents such as methylene chloride, sulfoxide-type solvents such as dimethyl

sulfoxide, amide-type solvents such as dimethylformamide, ether-type solvents such as tetrahydrofuran; dioxane and diglyme, alcohol-type solvents such as methanol and ethanol, nitrile-type solvents such as acetonitrile, ketone-type solvents such as acetone and cyclohexanone, ester-type solvents such as ethyl acetate, heterocyclic-type solvents such as pyridine and the like. Two or more kinds of organic solvents may be mixed and used.

For the reaction conditions, Miyaura, N., Suzuki, A., Chemical Review, 1995, vol. 95, p.2457; Snieckus, V., Chemical Review, 1990, vol. 90, p.879 and the like and references cited therein can be referred to.

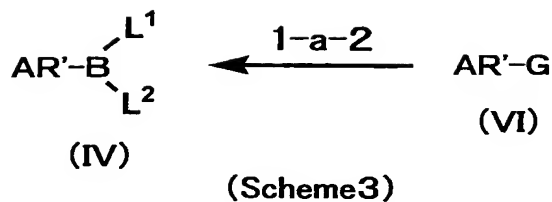
When hydroxyl group or amino group reactive under the aforementioned reaction conditions or inhibiting the reactions exists in the group AR', Rs' or V' in the aromatic ring (E'), this substituent is preferably protected.

When a protective group of hydroxyl group or amino group exist in the group AR', Rs' or V' in the aromatic ring (E') of the compound (Ia') prepared as described above, such a protective group can be eliminated during or after the preparation of Compound (Ia') to convert the compound into Compound (I) of the present invention. As for selection, introduction and deprotection of these protective groups of hydroxyl group and amino group, ordinary chemical publications, for example, Protective Groups In Organic Synthesis THIRD EDITION, John Wiley & Sons) and references cited therein can be referred to.

[Preparation Method 1] (Step a-2)

As Compound (IV), a compound commercially available as a reagent may be used, or as shown in the following scheme 3:

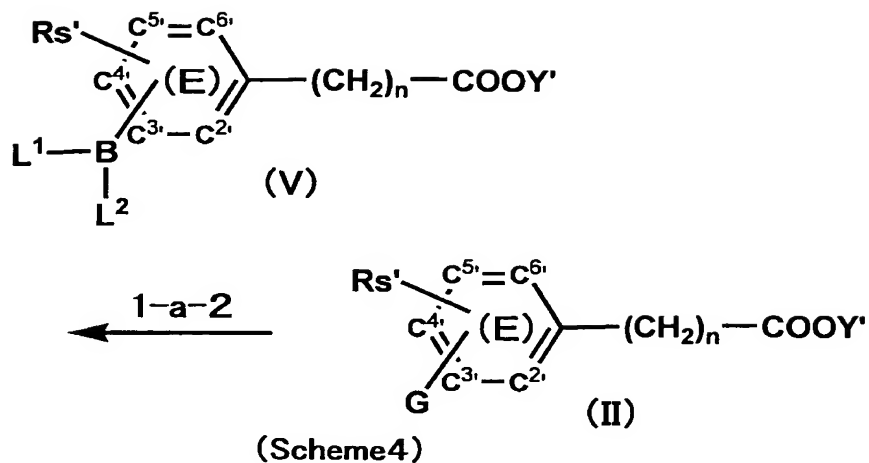




the compound can be produced from Compound (VI), which is commercially available or can be synthesized by a known method or a similar method thereto, according to the method described in the aforementioned reference (Chemical Review, vol. 95, p.2457, 1995) or the method described in Satoh, Y. et al., SYNTHESIS, p.1146, 1994 or according to the references cited therein.

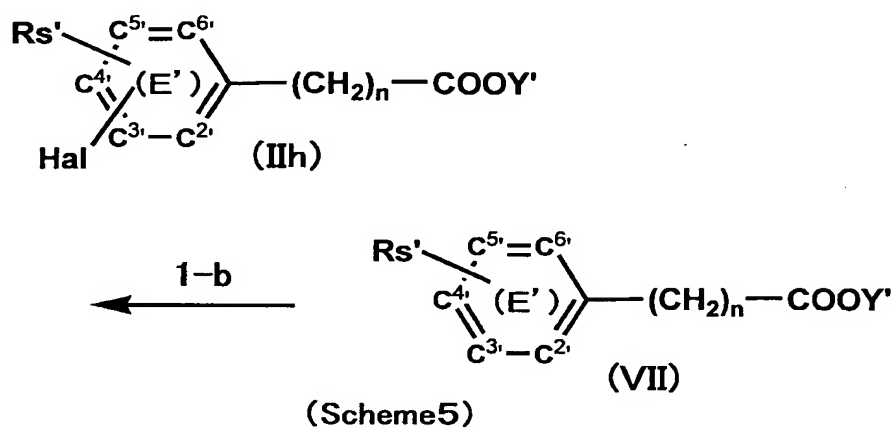
For example, examples include a method of preparing Compound (VI) by converting Compound (VI) into a lithio-compound using an alkyl lithium such as n-butyl lithium and t-butyl lithium, then reacting the product with a trialkyl borate and treating the product with a mineral acid such as hydrochloric acid, sulfuric acid, and phosphoric acid; and a method of preparing Compound (VI) by performing a cross-coupling reaction of Compound (VI) and an (alkoxy)diboron in the presence of a palladium catalyst and a base.

An example of the preparation method of Compound (V) includes a method of subjecting Compound (II) to a reaction similar to that of the aforementioned Step a-2, as shown in the following scheme 4:



[Preparation Method 1] (Step b)

As shown in the following scheme 5:

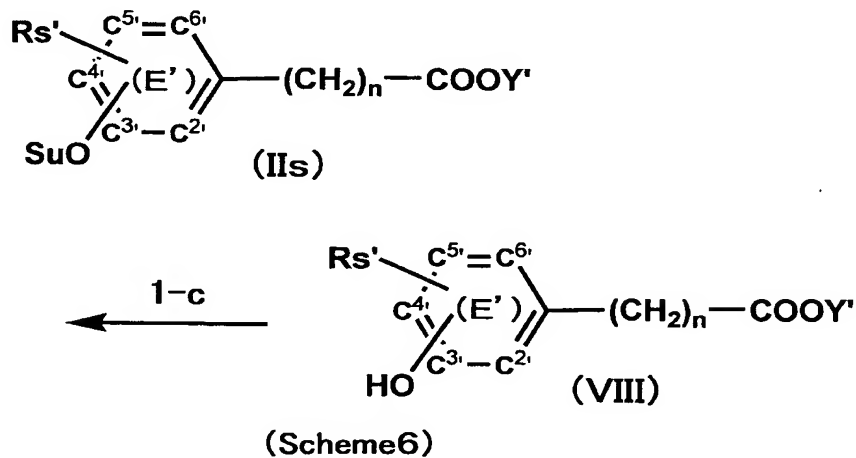


a compound represented by the formula (IIh) (hereinafter simply referred to as "Compound (IIh)"), which correspond to the compounds (II) wherein G represents a halogen atom such as chlorine atom, bromine atom or iodine atom, can be prepared by halogenating a compound represented by the formula (VII) [this compound is simply referred to as "Compound (VII)"], which is commercially available or can be prepared by a known method or a method similar thereto. In the formula of Compound (IIh), the group Hal represents a halogen atom, which may be any of

chlorine atom, bromine atom and iodine atom. As for the halogenation, examples of chlorination include a preparation method described in ordinary publications in the field of chemistry, for example, Shin Jikken Kagaku Koza (edited by Chemical Society of Japan, published by Maruzen Co., Ltd.), vol. 14, p.354. Examples of the method include a method utilizing chlorine (Cl<sub>2</sub>), a method utilizing sulfuric chloride, and the like. Examples of bromination include a preparation method described in ordinary publications in the field of chemistry, for example, Shin Jikken Kagaku Koza (edited by Chemical Society of Japan, published by Maruzen Co., Ltd.), vol. 14, p.354. Examples of the method include a method utilizing bromine (Br<sub>2</sub>), a method utilizing N-bromosuccinimide, and the like. Examples of iodination include a preparation method described in ordinary publications in the field of chemistry, for example, Shin Jikken Kagaku Koza (edited by Chemical Society of Japan, published by Maruzen Co., Ltd.), vol. 14, p.423. Examples of the method include a method utilizing iodine (I<sub>2</sub>), a method utilizing potassium triiodide, and the like.

[Preparation Method 1] (Step c)

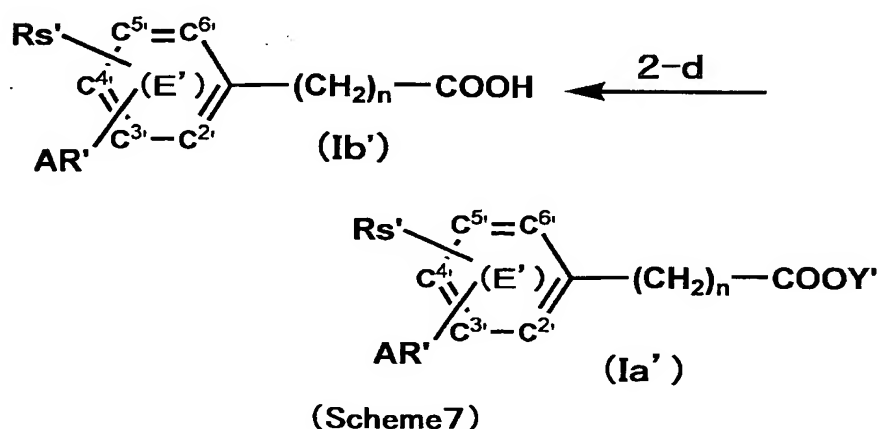
As shown in the following scheme 6:



a compound represented by the formula (II<sub>s</sub>) (this compound is hereinafter simply referred to as "Compound (II<sub>s</sub>)"), which corresponds to Compound (II) wherein G represents mesylate group, triflate group, or an arenesulfonate group, can be prepared by converting a compound represented by the formula (VIII) (this compound is simply referred to as "Compound (VIII)"), which is commercially available or can be prepared by a known method or a method similar thereto, into a sulfonic acid ester. In the formula of Compound (II<sub>s</sub>), the group Su represents methanesulfonyl group, trifluoromethanesulfonyl group, or arenesulfonyl group of which aromatic ring may be substituted with one of T<sup>1</sup> or two or more of identical or different T<sup>1</sup>. Examples of the method for the conversion into sulfonic acid ester include a preparation method described in ordinary publications in the filed of chemistry, for example, Shin Jikken Kagaku Koza (edited by Chemical Society of Japan, published by Maruzen Co., Ltd.), vol. 14, p.1793. Examples of the method include a method utilizing sulfonyl chloride, a method utilizing sulfonic anhydride, and the like.

[Preparation Method 2] (Step d)

As shown in the following scheme 7:



a compound represented by the formula (Ib') wherein Y represents hydrogen atom,

and Rs, AR, and V on or in the aromatic ring (E) may be protected (this compound is hereinafter simply referred to as "Compound (Ib')"), which constitutes a part of the scope of Compound (I) of the present invention, can be prepared by hydrolyzing Compound (Ia') so as to convert the group OY' into hydroxyl group.

For the reaction of converting Compound (Ia') into Compound (Ib'), in general, the compound is preferably reacted in a base. Further, for the reaction of converting Compound (Ia') to Compound (Ib'), in general, the compound is preferably reacted in an inert medium that does not inhibit the reaction, preferably a polar solvent.

Examples of the base used in the above reaction include, for example, alkali metal bases such as sodium hydroxide, potassium hydroxide, sodium carbonate, potassium carbonate, sodium methoxide and potassium t-butoxide and organic bases such as triethylamine. As for amounts of the bases, generally 1 to 20 moles, preferably 1 to 10 moles, for alkali metal bases, or 1 to a large excess moles for organic bases based on Compound (Ia').

Examples of the polar solvent include water, methanol, ethanol, tetrahydrofuran, dioxane and the like, and these solvents may be used as a mixture as required. As the reaction temperature, an appropriate temperature of, for example, from room temperature to a refluxing temperature of solvent is chosen. The reaction time is, for example, generally 0.5 to 72 hours, preferably 1 to 48 hours, when an alkali metal base is used, or generally 5 hours to 14 days when an organic base is used. Since progress of the reaction can be monitored by thin layer chromatography (TLC), high performance liquid chromatography (HPLC) or the like, the reaction can generally be terminated appropriately so as to maximize the yield of Compound (Ib').

For collection of Compound (Ib') obtained as described above from the reaction solution as a free carboxylic acid, operations may preferably be carried out

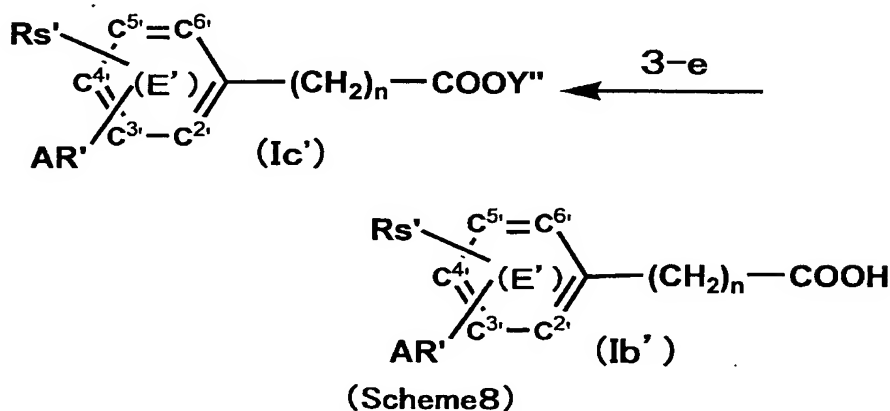
by, when the polar solvent is a water-soluble solvent, evaporating the solvent, neutralizing the residue with an inorganic acid such as aqueous hydrochloric acid, dissolving the residue in a water-insoluble solvent, then washing the solution with a weakly acidic aqueous solution, water or the like, and evaporating the solvent. When the polar solvent is a water-insoluble solvent, operations may preferably be carried out by neutralizing the reaction solution with an inorganic acid, washing the solution with a weakly acidic aqueous solution, water or the like, and then evaporating the solvent.

Further, when Compound (Ib') forms a salt with the base used after the reaction to give a solid, the salt of Compound (Ib') can be obtained by isolation and purification of the solid in a conventional manner.

When a protective group of hydroxyl group or amino group exists in the group AR', Rs' or V' in the aromatic ring (E') of Compound (Ia') prepared as described above, Compound (Ia') can be converted into Compound (I) of the present invention by removing the protective group during or after the preparation of Compound (Ia').

[Preparation Method 3] (Step e)

As shown by the following scheme 8:



a compound represented by the formula (Ic') [hereinafter simply referred to as "Compound (Ic')"] as Compound (I) of the present invention wherein the group Y represents Y", and Rs, AR, and V in the aromatic ring (E) may be protected, can be produced by esterifying the carboxyl group (COOH) of Compound (Ib') in a conventional manner. In the formula of Compound (Ib'), Y" represents a lower alkyl group having 1 to 4 carbon atoms, a  $-(CH_2)_mNR^{18}R^{19}$  group, or  $C(R^{20})_2OC(O)A^3R^{21}$ .

Examples of the method for producing Compound (Ic') include a method of allowing Compound (Ib') to react with an inorganic halide without solvent or in an inert solvent to convert the compound into an acid halide and then allowing the acid halide per se or the same dissolved in an inert solvent to react with an excess amount of hydroxide of the targeted Y". Examples of the inorganic halide used in this method include thionyl chloride, phosphoryl chloride, phosphorus pentachloride, phosphorus trichloride and the like, and thionyl chloride is a preferred example. Examples of an amount used include generally an equimolar to a large excess amount, preferably 1.5 to 5 moles based on Compound (Ib'). Examples of the inert solvent used in this reaction include, for example, halogenated hydrocarbons such as dichloromethane, chloroform and 1,2-dichloroethane, ethers such as tetrahydrofuran and dioxane, and benzene compounds such as benzene, toluene, xylene and chlorobenzene. These solvents can be used, for example, each alone or as a mixed solvent. In order to promote the reaction, a catalytic amount of N,N-dimethylformamide may be added. As a reaction temperature, an appropriate temperature of from room temperature to a refluxing temperature of the solvent is generally chosen. Examples of the reaction time include generally 0.5 to 24 hours, preferably 1 to 6 hours.

Examples of the inert solvent used for the reaction with hydroxide of the targeted Y" include, for example, halogenated hydrocarbons such as

dichloromethane, chloroform and 1,2-dichloroethane, ethers such as tetrahydrofuran and dioxane, and benzene compounds such as benzene, toluene, and xylene. The reaction can also be performed with an excess amount of the hydroxide of the targeted Y" without using a solvent. As the reaction temperature, an appropriate temperature of from -10°C to room temperature is chosen. Examples of the reaction time include generally 0.5 to 24 hours, preferably 0.5 to 6 hours.

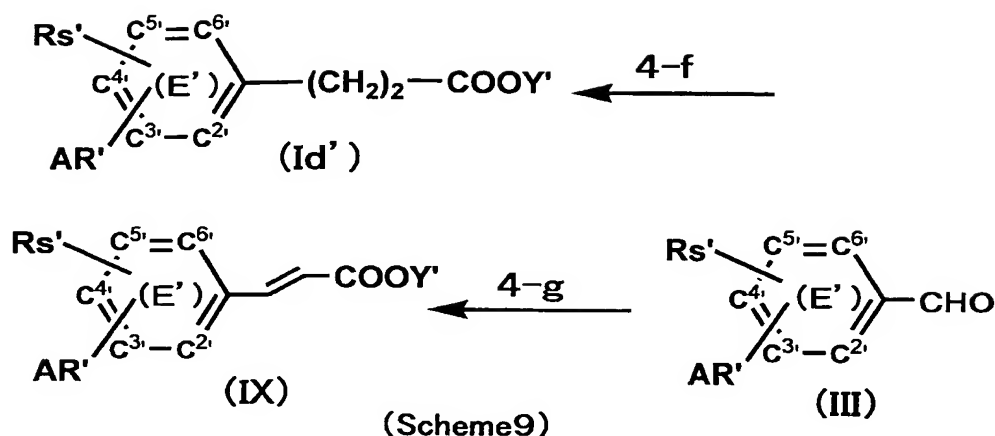
Other methods for producing Compound (Ic') include, for example, the "esterification using an alcohol" described in Shin Jikken Kagaku Koza (edited by the Chemical Society of Japan, published by Maruzen Co., Ltd.), vol. 14, p.1002, "esterification using an O-alkylating agent", ibid, the same volume, p.1002, "esterification using an alkyl halide", ibid, the same volume, p.1008, "esterification reaction using a dehydrating agent", ibid, vol. 22, p.45 and the like.

When hydroxyl group or amino group reactive under the aforementioned reaction conditions or inhibiting the reactions exists in AR', Rs' or V' in the aromatic ring (E'), this substituent is preferably protected.

When a protective group of hydroxyl group or amino group exist in AR', Rs' or V' in the aromatic ring (E') of the compound (Ic') prepared as described above, such a protecting group can be eliminated during or after the preparation of Compound (Ic') to convert the compound into Compound (I) of the present invention. [Preparation Method 4]

As shown in the following scheme 9:





a compound represented by the formula (Id') (hereinafter this compound is simply referred to as "Compound (Id')") as Compound (I) of the present invention wherein n in the methylene moiety is an integer of 2, and wherein Rs, AR, and V in the aromatic ring (E) may be protected, can also be prepared by the method shown below.

[Preparation Method 4] (Step f)

Compound (Id') can be prepared by reducing the double bond of a compound represented by the formula (IX) (hereinafter this compound is simply referred to as "Compound (IX)") using a reduction reaction described in ordinary publications in the field of chemistry. Examples of the reaction include a method of converting the double bond of Compound (IX) into a single bond by hydrogenation using a hydrogen source such as hydrogen gas, ammonium formate, and hydrazine hydrate in a single solvent or a mixed solvent of alcoholic-type solvents such as methanol, ester-type solvents such as ethyl acetate in the presence of a catalyst such as palladium/carbon powder, platinum oxide (PtO<sub>2</sub>), and activated nickel.

When hydroxyl group or amino group reactive under the aforementioned reaction conditions or inhibiting the reactions exists in AR', Rs' or V' in the aromatic ring (E'), this substituent is preferably protected.

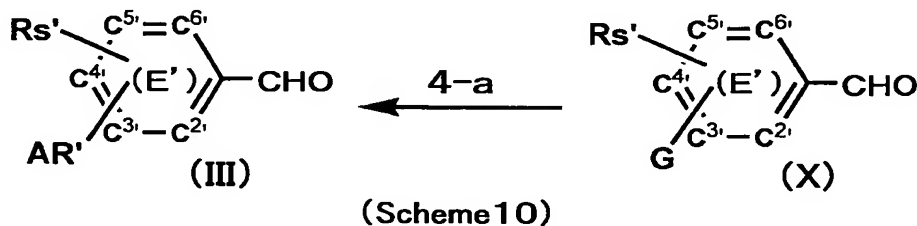
When a protective group of hydroxyl group or amino group exist in AR', Rs'

or V' in the aromatic ring (E') of the compound (Id') prepared as described above, such a protecting group can be eliminated during or after the preparation of Compound (Id') to convert the compound into Compound (I) of the present invention. [Preparation Method 4] (Step g)

Compound (IX) can be prepared from a compound represented by the formula (III) [hereinafter this compound is simply referred to as "Compound (III)"]. Examples of the preparation method include a method utilizing the Horner-Emonds reaction described in Shin Jikken Kagaku Koza (edited by Chemical Society of Japan, published by Maruzen Co., Ltd.), vol. 14, p.238. Specifically, the compound can be obtained by reacting Compound (III) with a commercially available dialkylphosphonoacetic acid ester in an inert solvent, for example, an alcohol-type solvent such as methanol and ethanol or ether-type solvent such as tetrahydrofuran and dimethoxyethane in the presence of a base such as sodium hydride and sodium alkoxide. As the reaction temperature, an appropriate temperature of from -10°C to a refluxing temperature of a solvent is generally chosen, and preferred examples include a temperature of from 0°C to room temperature. The reaction time is generally 1 to 16 hours, preferably 2 to 8 hours. Since progress of the reaction can be monitored by thin layer chromatography (TLC), high performance liquid chromatography (HPLC) or the like, the reaction can generally be terminated appropriately so as to maximize the yield of Compound (IX).

[Preparation Method 4] (Step a)

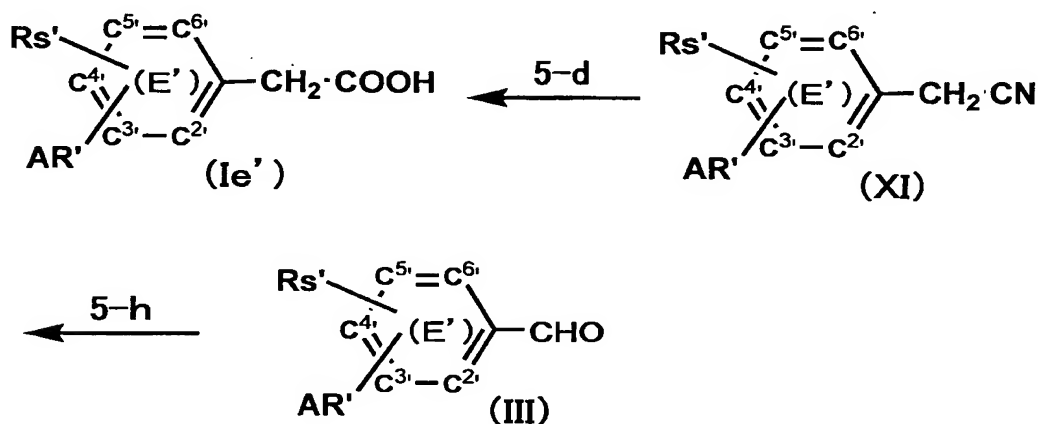
As shown in the following scheme 10:



Compound (III) can be prepared by introducing the substituent AR' into a compound represented by the formula (X) [hereinafter this compound is simply referred to as "Compound (X)"] according to any of the methods described in the step a-1 of the preparation method 1 mentioned above.

[Preparation Method 5]

As shown in the following scheme 11:



(Scheme 11)

a compound represented by the formula (Ie') [hereinafter this compound is simply referred to as "Compound (Ie')"], as Compound (I) of the present invention wherein n in the methylene moiety is an integer of 1, Y represents hydrogen atom, and Rs, AR, and V in the aromatic ring (E) may be protected, can also be prepared by the method shown below.

[Preparation Method 7] (Step d)

Specifically, Compound (Ie') can be prepared by hydrolyzing nitrile group of a compound represented by the formula (XI) [hereinafter this compound is simply referred to as "Compound (XI)"] into carboxyl group according to a method similar to the method shown in the step d of the preparation method 2 mentioned above.

When a protective group of hydroxyl group or amino group exist in AR', Rs' or V' in the aromatic ring (E') of the compound (Ie') prepared as described above, such a protecting group can be eliminated during or after the preparation of Compound (Ie') to convert the compound into Compound (I) of the present invention. [Preparation Method 5] (Step h)

Compound (XI) can be produced from Compound (III) mentioned above. For example, Compound (III) is reacted with a trimethylsilyl cyanide using a Lewis acid, particularly zinc iodide, as a catalyst in an inert solvent such as tetrahydrofuran as described in Jikken Kagaku Koza, 4th Edition (edited by Chemical Society of Japan, published by Maruzen Co., Ltd.), vol. 20, p.445. Then, the reduction reaction using a hydrosilane described in Jikken Kagaku Koza, 4th Edition (edited by Chemical Society of Japan, published by Maruzen Co., Ltd.), vol. 26, p.197 is performed. Examples of the method of the reduction reaction include a method of performing the reduction with a hydrosilane such as triethylsilane and a protonic acid such as trifluoroacetic acid or a Lewis acid such as boron trifluoride in a halogenated solvent such as dichloromethane.

The preparation method of Compound (I) is not limited to the methods described herein. For example, the compounds of the present invention can be produced by modifying or converting a substituent of a compound serving as a precursor of the compounds according to a method or a combination of methods described in ordinary publications in the filed of chemistry.

Examples of the preparation method for Compound (I) of the present invention which contains an asymmetric carbon in the substituent Rs include a method of using a starting material in which a moiety corresponding to the asymmetric carbon in the substituent Rs is already optically active, which is commercially available (or can be prepared by a known method or a method similar thereto). A method is also available in which the compound of the present

invention or a precursor thereof is separated as an optically active isomer in a conventional manner. Examples of such method include, for example, a method utilizing high performance liquid chromatography (HPLC) using a chiral column, a method comprising condensation with an optically active reagent to form a diastereomer, successive separation and purification, followed by decomposition. When a precursor is separated to obtain an optical isomer, optically active Compound (I) of the present invention can then be prepared by performing the aforementioned preparation methods.

When Compound (I) of the present invention contains an acidic functional group such as carboxyl group or phenolic hydroxyl group, the compound can be converted into pharmaceutically acceptable salt (e.g., inorganic salts with sodium, ammonia and the like, or organic salts with triethylamine and the like) by a known means. For example, when an inorganic salt is to be obtained, it is preferable to dissolve Compound (I) of the present invention in water containing at least 1 equivalence of hydroxide, carbonate, bicarbonate or the like corresponding to a desired inorganic salt. For the reaction, an inactive water-miscible organic solvent such as methanol, ethanol, acetone, and dioxane may be mixed. For example, by using sodium hydroxide, sodium carbonate, or sodium hydrogencarbonate, a solution of sodium salt can be obtained.

When Compound (I) of the present invention contains a basic functional group such as amino group, or when Compound (I) of the present invention contains an aromatic ring which itself has properties of base (e.g., pyridine ring), the compound can be converted into a pharmaceutically acceptable salt (e.g., salt with inorganic acids such as hydrochloric acid and sulfuric acid, or salts with organic acids such as acetic acid and citric acid) by a known means. For example, when an inorganic salt is to be obtained, it is preferable to dissolve Compound (I) of the present invention in water containing at least 1 equivalence of a desired inorganic

acid. For the reaction, an inactive water-miscible organic solvent such as methanol, ethanol, acetone, and dioxane may be mixed. For example, by using hydrochloric acid, a solution of hydrochloride can be obtained.

When a solid salt is desired, a solution may be evaporated, or a water-miscible organic solvent having polarity to some extent, such as butanol or ethyl methyl ketone, can be added to obtain a solid salt thereof.

The various compounds disclosed by the present invention can be purified by known methods such as recrystallization, and variety of chromatography techniques (column chromatography, flash column chromatography, thin layer chromatography, high performance liquid chromatography).

Compound (I) of the present invention and pharmaceutically acceptable salts thereof have an action of suppressing the production of both of prostaglandins and leukotrienes. The action of suppressing the production of prostaglandins and/or leukotrienes includes, for example, an action of suppressing PGE<sub>2</sub> production, observed when cultured cells of MG-63 which is a human osteosarcoma cell line are stimulated with IL-1  $\beta$  and/or PGD<sub>2</sub> and LTB<sub>4</sub> production observed when cultured cells of RBL-2H3 which is a rat mastocytoma cell line are stimulated with IgE, by 10% or more, preferably 30% or more, most preferably 50% or more, compared with a positive control at a concentration of the compound not having cytotoxicity. As for a mode of action at a molecular level, it is considered that the compound of the present invention inhibits both of COX-1 and/or COX-2, which produce prostaglandins, and 5-LO, which produces leukotrienes. It is also considered that the compound of the present invention suppresses the production of arachidonic acid by inhibiting enzymatic activity of type 2A, 4, or 5 PLA<sub>2</sub> involved in prostaglandin and leukotrien production.

It is considered that, in these molecular action mechanisms, Compound (I) of the present invention inhibits the enzymatic activity of type 4 PLA<sub>2</sub>. For the

judgment, for example, the enzyme inhibitory action against type 4 PLA<sub>2</sub> can be examined, and known methods for measuring the enzymatic activity of type 4 PLA<sub>2</sub> are preferably utilized [Clark et al., Proceeding of National Academy of Science USA (Proc. Natl. Acad. Sci. USA), 1990, vol. 87, p.7708; Gronich et al., Biochemical Journal (Biochem. J.), 1990, vol. 271, p.37; Clark et al., Cell, 1991, vol. 65, p.1043; Kramer et al., Journal of Biological Chemistry (J. Biol. Chem), 1991, vol. 266, p.5268]. The type 4 PLA<sub>2</sub> inhibitory action of the compounds of the present invention can be elucidated by employing these methods.

Compounds (I) of the present invention and pharmaceutically acceptable salts thereof inhibited mouse inflammatory edema, allergic edema, acetic acid writhing reaction, and rat adjuvant arthritis by oral administration at a dose of 0.1 to 500 mg/kg, and caused no death of the mice by oral administration at a dose of 500 mg/kg/day for 3 days. Therefore, they are safe compounds as drugs for mammals, preferably humans, pets or companion animals such as dogs and cats, and farm animals, and they are useful substances as active ingredients of medicaments. Preferred examples of the medicaments for mammals, preferably humans, pets or companion animals such as dogs and cats, and farm animals include agents for prophylactic and/or therapeutic treatment of various conditions, various diseases, and pathological conditions in which an acute or chronic inflammatory reaction resulted from production of prostaglandin and/or leukotriene is observed, specifically inflammatory diseases, allergic diseases, autoimmune diseases, and pain.

More specifically, the conditions or diseases include arthritis, chronic rheumatoid arthritis, malignant rheumatoid arthritis, juvenile rheumatoid arthritis, Felty's syndrome, adult Still's disease, osteoarthritis, synovitis, gout, slack of artificial joint implant, fervescence, common cold, algesia, burn, thermal injury, keloplasty, menstrual pain, dysmenorrhea, menstrual cramp, allergic

reaction, allergic contact hypersensitivity, allergic rhinitis, pollinosis, allergic conjunctivitis, hypersensitivity pneumonitis, allergic bronchopulmonary mycosis, emphysema, acute respiratory distress syndrome, asthma, bronchitis, chronic obstructive pulmonary disease, chronic bronchitis, pulmonary emphysema, diffuse panbronchiolitis, respiratory obstruction, graft versus host syndrome, urticaria, ultraviolet radiation dermatitis, atopic dermatitis, cancer, myelogenous leukemia, sarcomata, brain tumor, cachexia, tissue ulcer, digestive ulcer, gastritis, acute and chronic pancreatitis, regional enteritis, ulcerative colitis, diverticulitis, recurrent gastroenteric disorder, gastroenteric bleeding, inflammatory bowel disease, Crohn's disease, intestinal tract type Behcet's disease, infectious enteritis, ischemic enteritis, radiation enteritis, drug-induced enteritis, irritable bowel syndrome, hepatic diseases (hepatopathies, liver failures) such as acute hepatitis, fulminant hepatitis, chronic hepatitis, hepatic cirrhosis, fatty liver, alcoholic liver injury, drug liver injury (drug-induced hepatitis), congestive hepatitis, autoimmune hepatitis, primary biliary cirrhosis and hepatic porphyria, coagulation, anemia, ankylosing spondylitis, restenosis, periodontosis, epidermolysis bullosa, atherosclerosis, aortic aneurysm, periarteritis nodosa, congestive cardiac failure, arrhythmia, myocardial infarction, cerebral infarction, attack, cerebral ischemia, head injury, spinal cord injury, myelopathic muscular atrophy, neuralgia, neurodegenerative disease, Alzheimer's disease, Lewy body disease, Shy-Drager syndrome, Reye's syndrome, progressive supranuclear palsy, progressive multifocal leukoencephalopathy, normal pressure hydrocephalus, subacute sclerosing panencephalitis, frontal lobe type dementia, acute anterior poliomyelitis (poliomyelitis), poliomyelitis neurosis, viral encephalitis, Creutzfeldt-Jakob disease, Kuru disease, bovine spongiform encephalopathy (mad cow disease), scrapie, epilepsy, cerebral amyloid angiopathy, autoimmune disease, Huntington's disease, Parkinson's disease, migraine, depression, mania, manic-depressive psychosis, hereditary cerebellar ataxia,



peripheral neuropathy, glaucoma, pain, gingivitis, postoperative pain, amyotrophic lateral sclerosis, osteoporosis, multiple sclerosis, ocular angiogenesis, cornea damage, macular degeneration, conjunctivitis, abnormal wound healing, sprain or strain of muscle or joint, tendinitis, skin disease, psoriasis vulgaris, pustular psoriasis, erythroderma psoriaticum, arthritic psoriasis, myasthenia gravis, multiple myositis, myositis, bursitis, diabetes mellitus, tumor invasion, tumor growth, tumor metastasis, cornea scar, scleritis, immunodeficiency disease, pachydermia, eosinophilic fasciitis, sepsis, endotoxin shock, premature delivery, hypoprothrombinemia, hemophilia, thyroiditis, sarcoidosis, Behcet's syndrome, hypersensitivity, renal disease, rickettsial infectious disease, protozoal disease, reproduction disease, sepsis shock and the like. Other specific conditions and diseases include toothache, pain after tooth extraction, back or low back pain, periarthrititis humeroscapularis, cervico-omo-brachial syndrome, tenosynovitis, acute upper respiratory inflammation, herpes zoster, fibrosis, pulmonary fibrosis, pneumoconiosis, chronic interstitial pneumonia, granulomatous interstitial pneumonia, fibrosing interstitial pneumonia, renal fibrosis, nephropylitis, various types of secondary contracted kidney, glomerular nephritis, chronic nephritis, glomerulosclerosis, hepatic fibrosis, cardiac fibrosis after myocardial infarction, idiopathic cardiomyopathy, pancreatic sclerosis, pancreatic fibrosis, pancreatolithiasis, Takayasu's arteritis, chronic thyroiditis, dermatomyositis, multiple myositis, myelofibrosis, Banti disease, retroperitoneal fibrosis, various radiation injuries and the like. Further, the medicament comprising Compound (I) of the present invention as an active ingredient can be used for the aforementioned conditions or diseases of mammals, preferably humans, pets or companion animals such as dogs and cats or farm animals together with or in combination with one or more kinds of other prophylactic or therapeutic drugs.

Examples of the drugs that can be used together or in combination include,

for example, the following drugs: immunomodulation-type antirheumatic drugs and antimetabolite used as therapeutic drugs for rheumatoid arthritis, specifically, gold preparations, bucillamine, lobenzarit, salazosulfapyridine, methotrexate, azathiopurin, mizoribine, leflunomide, tacrolimus, cyclosporin and the like and preparations containing the same; anti-cytokine antibody preparations directed to cytokines such as interleukin (IL) 1, IL-6, and tumor necrosis factor (TNF)- $\alpha$  or preparations of soluble receptors for those cytokines, which are biological preparations, specifically, infliximab, etanercept and the like and preparations containing the same; steroid preparations, specifically, dexamethasone, betamethasone, prednisolone, fluticasone, beclometasone and the like and preparations containing the same; bronchodilators used as therapeutic agents for chronic bronchial asthma, specifically, salmeterol and salbutamol, which are adrenalin  $\beta$  2 stimulants, ipratropium, which is an anticholinergic drug, and the like and preparations containing the same; therapeutic drugs for allergic diseases, for example, theophylline, which is a xanthine analogue drug, and the like, fexofenadine, epinastatine, cetirizine, ketotifen, disodium cromoglycate, pemirolast and the like, which are antiallergic agents, fexofenadine, cetirizine and the like, which are antihistaminic agents, and preparations containing the same; irinotecan, 5-fluorouracil and the like, which are antitumor agents, and preparations containing the same. Further, the medicament comprising Compound (I) of the present invention as an active ingredient is used, for example, together with or in combination with radiotherapy.

In order to use Compound (I) of the present invention or pharmaceutically acceptable salts thereof for the medicaments described above, an effective amount of Compound (I) of the present invention or a pharmaceutically acceptable salt thereof, per se, may be used, or the substance may be mixed with a pharmaceutically acceptable carrier to form a pharmaceutical composition. The

carrier may be, for example, a suspending agent such as carboxymethylcellulose, or purified water, physiological saline or the like, if desired. Other known carriers can also be used. Examples include a method of dissolving Compound (I) of the present invention or a pharmaceutically acceptable salt thereof in purified water containing 0.5% carboxymethylcellulose and using the solution.

Examples of formulations for preparing the aforementioned pharmaceutical composition include tablet, powder, granule, syrup, suspension, capsule, and injection. For the manufacture of these formulations, various carriers suitable for these preparations are used. For example, examples of the carrier for oral preparations include excipients, binders, lubricants, fluid accelerators, and colorants.

When the compound of the present invention is formulated as a parenteral preparation such as an injection, water for injection, physiological saline, glucose aqueous solution, vegetable oil for injection, propylene glycol, polyethylene glycol and the like can generally be used as a diluent. Disinfectants, antiseptics, stabilizers, isotonic agents, soothing agents and the like may be further added, as required.

When the compound of the present invention is administered to a mammal, e.g., human, the compound can be administered in the form of a tablet, a powder, a granule, a suspension, a capsule or the like. The compound can also be parenterally administered in the form of a suppository, a gel, a lotion, an ointment, a cream, or a spray. A dose thereof varies depending on a disease to be applied, administration route, age, weight, degree of symptom of a patient and the like. Examples of the dose include generally an administration at a dose of 1 to 1,000 mg per day for an adult once to three times a day. Every day administration for a period of several days to two months is commonly applied. The daily dose and the administration period may be increased or decreased depending on symptoms of a

patient.

Fibrosis, which is a disease characterized by fibrosing of tissues, is known as a severe disease which is often mortal. Fibrosing of tissues is caused by proliferation of interstitial cells, which represented by fibroblasts, and production of extracellular matrix such as collagen. Fibrosing is considered a repair mechanism against tissue affections in organs. Excessive fibrosing causes fibrosing diseases of organs, and further progression of fibrosing causes sclerotic diseases. Many of such sclerotic diseases are intractable, progressive and irreversible. Although fibrosing varies in various organs, etiological hypotheses of fibrosing have many similarities. That is, a certain inflammatory lesion precedes, and in its healing process, various kinds of cytokines and growth factors are produced mainly from immunocompetent cells and platelets as well as interstitial cells such as fibroblasts themselves involved in the healing, and activated to cause deposition of extracellular matrix (Takehara, Molecular Medicine, 2001, vol. 38, p.854).

Among fibroses, pulmonary fibrosis is one of the representative diseases. Pulmonary fibrosis is a disease in which disruption of alveolar structure is caused by chronic inflammation and increase of collagenic fibers in alveolar walls, and which eventually leads to respiratory failure and death. For example, pulmonary fibrosis occurs following infectious pneumonia and the like. Examples of the infectious pneumonia include severe acute respiratory syndrome (SARS) and influenzal pneumonia. It has been reported that, in SARS, in particular, severe inflammation is caused in pulmonary stroma, and as a result, it highly likely to develop into pulmonary fibrosis (Antonino et al., Radiology, 2003). In addition, pulmonary fibrosis is also caused by various medicaments.

In recent years, with increase of medicaments used for diagnosis, prophylactic and therapeutic treatments of various kinds of diseases, drug-induced

pulmonary fibrosis caused by such drugs is increasing. Drug-induced pulmonary fibrosis is a severe disease that eventually leads to death, and it causes serious problems in therapeutic treatments of various diseases. Therefore, prophylactic and therapeutic treatments of drug-induced pulmonary fibrosis constitute a particularly important subject of concern.

Against drug-induced pulmonary fibrosis, steroid therapy is currently used. However, effective rate of the steroid therapy is low and the effect is only partial and transient, and thus lesions often remain [Igaku no Ayumi, 2001, vol. 197, p.313]. Further, side effect of steroid agents and acute aggravation due to decrease of doses or termination of their administrations are also often observed, which remains clinically far unsatisfactory level.

As a recent finding, it was reported that administration of pirfenidone was effective against pulmonary fibrosis in clinical tests in the United States (Raghu et al., American Journal of Respiratory and Critical Care Medicine, 1999, vol. 159, p.1061) and Japan (Nagai et al., Internal Medicine, 2002, vol. 41, p.1118). However, development of novel prophylactic and/or therapeutic agents highly effective for these diseases is desired at all events.

The medicament provided by the present invention is useful as a medicament containing a type 4 PLA2 inhibitor as an active ingredient for prophylactic and/or therapeutic treatment of fibrosis, preferably pulmonary fibrosis, further preferably drug-induced pulmonary fibrosis.

As described above, fibrosis, in particular, pulmonary fibrosis, is a severe disease and is an important object of prophylactic and/or therapeutic treatment. As for pulmonary fibrosis, more than 100 kinds of factors including toxic gases and various medicaments have been elucidated as the causes of early alveolopathy. As described above, with the increase of medicaments used for diagnosis, prophylactic and therapeutic treatments of various kinds of diseases, drug-induced pulmonary

fibrosis caused by such drugs is increasing.

As for drug-induced pulmonary fibrosis, causality between expression of pathological conditions such as coughing, difficulty of breathing, or fervescence and the administration of medicaments is suspected, and it is considered that a diffuse interstitial shadow appears on a thoracic X-ray photograph simultaneously with or slightly after the administration of medicaments.

As medicaments reported to cause drug-induced pulmonary fibrosis, anticancer agents, anti-rheumatic agents, immunosuppressants, antibiotics, chemotherapeutants, antihypertensive agents, diuretics, anti-inflammatory/analgesic agents, biologics, Chinese medicines are known (Inooka et al., Therapeutics, 1995, vol. 29, p.1295). Typical medicaments are shown in Table 1.

Table 1

Classification	Examples of agent
1) Anticancer agent, immunosuppressant	Peplomycin, bleomycin, cyclophosphamide, nitrosourea, busulfan, methotrexate, azathioprine, mitomycin-C, tegafur, carmofur, tegafur/uracil preparation, cisplatin, doxorubicin, 6-mercaptopurine, daunomycin, vincristine, vinblastine, vindesine, procarbazine, neocarzinostatin, melphalan, thiotepa, nimustine, cytarabine, zinostatin, stimalamer, chlorambucil, carmustine, lomustine, semustine, teniposide, etoposide, Taxol, taxotere, irinotecan, gefitinib, tamoxifen and the like.
2) Antihypertensive agent, diuretic	$\alpha$ -methyldopa, trichlormethiazide, hydrochlorothiazide, enalapril, hexamethonium, mecamlamine, pentolinium,

	practolol, pindolol, propranolol, acebutolol, hydralazine
3) Antibiotic, chemotherapeutant	Cephem antibiotics (cephaloridine, cephalothin, cephalixin, cefradine, cefazolin, cefaclor, cefmenoxime, cefmetazole, cefoperazone, cefotiam, cefroxadin, ceftizoxime, latamoxef and the like), tetracyclines (minocycline, oxytetracycline), antituberculous agents (isoniazid, paraaminosalicylic acid, rifampicin, streptomycin), penicillin antibiotics (ampicillin, piperacillin, nafcillin, penicillin, amoxicillin), aminoglycoside antibiotics (streptomycin), macrolide antibiotics (erythromycin), phosphomycin, aminoglycosides (tobramycin, Micromycin), new quinolone drugs (enoxacin, ofloxacin, norfloxacin), antifungal agents (amphotericin) and the like
4) Others	Inhalants (cromoglicic acid and the like), gold preparations (aurothiomalic acid and the like), psychotropic agents and nervines (amitriptyline, diphenylhydantoin, carbamazepine, phenobarbital, valproate salt, imipramine, mephensin, meproamate), antiphlogistic and analgesics (naproxen, acetaminophen, acetylsalicylic acid, phenacetin, diclofenac, loxoprofen, fenbufen, nabumetone, aluminoprophen and the like), antiarrhythmic agents (amiodarone, procainamide, aprindine), antidiabetic agents (chlorpropamide), antithyroid agents (thiouracil), proteolytic enzymes (serrapeptidase), antiparkinsonic agents (levodopa,

	bromocriptine), antirheumatic agents (bucillamine, auranofin, actarit), sho-saiko-to, chai-ling-tang, rikkunshi-to, interferon, warfarin, salazosulfapyridine, dichloroferamide, fominoben, D-penicillamine, propylthiouracil, corticosteroid, flavoxate, allopurinol, ethoxysclerol and the like
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In therapeutic treatment of rheumatoid arthritis, for example, agents that cause pulmonary fibrosis at high frequency such as methotrexate and sodium aurothiomalate are used as disease-modifying antirheumatic drugs. Further, disease-modifying antirheumatic drugs that may cause pulmonary fibrosis at a relatively low frequency, such as actarit, bucillamine, auranofin, salazosulfapyridine, and D-penicillamine are also used. Although these disease-modifying antirheumatic drugs are useful agent in the rheumatoid arthritis treatment system, pulmonary fibrosis caused as a side effect is a factor of restricting use of these drugs. In recent years, methotrexate, in particular, has come to be used as an antirheumatic agent, and onset of pulmonary fibrosis that is also histopathologically called interstitial pneumonia as the side effect of methotrexate becomes a problem in the rheumatoid arthritis treatment system.

Further, in cancer therapy, cyclophosphamide, Taxol, etoposide, cisplatin, vincristine, vinblastine, irinotecan, gefitinib, and bleomycin are useful as anticancer agents. However, because all of these anticancer agents cause pulmonary fibrosis that is also histopathologically called as interstitial pneumonia as a side effect at a high frequency, they have a problem in the therapeutic treatment system. Bleomycin, gefitinib, irinotecan, and cisplatin are used for therapeutic treatment of lung cancer. However, if patients with lung cancer develop pulmonary fibrosis, the condition is most likely for the patients to be fatal.



Among these drugs, bleomycin suffers from a problem that it causes pulmonary fibrosis at a high frequency.

Preferred objects of application of the medicament of present invention are drug-induced pulmonary fibroses caused by these drugs.

In present invention, the type 4 PLA<sub>2</sub> inhibitor is not particularly limited so long as the inhibitor has type 4 PLA<sub>2</sub> inhibitory activity. For example, known type 4 PLA<sub>2</sub> inhibitors can be chosen. Examples of the known type 4 PLA<sub>2</sub> inhibitors include the following inhibitors: the compounds described in U.S. Patent No. 5,462,954, preferably 2-phenyl-4-ethyl-5-[6-(2H-tetrazol-5-yl)-6-methylheptyloxy]phenol, 8-propyl-7-{3-[4-(4-fluorophenyl)-2-ethyl-5-hydroxyphenoxy]propyloxy}-3,4-dihydro-2H-1-benzopyran-2-carboxylic acid, and 2-{3-[3-([5-ethyl-2-hydroxy(1,1'-biphenyl)-4-yl]oxy)propyloxy]-2-propylphenoxy}propionic acid; the compounds described in WO99/43654, preferably 4-(1-benzhydryl-6-chloro-1H-indol-3-ylmethyl)-3-methoxybenzoic acid; the compounds described in WO98/33797, preferably N-{4-(biphenyl-2-ylmethyl-isobutylamino)-1-[2-(4-fluorobenzoyl)benzoyl]pyrrolidin-2-ylmethyl}-3-[4-(2,4-dioxothiazolidin-5-ylidenemethyl) phenyl]acrylamide and the like; the compounds described in WO01/30387, preferably N-{1-[2-(2,4-difluorobenzoyl)benzoyl]-4-tritylsulfanylpyrrolidin-2-ylmethyl}-4-(2,4-dioxothiazolidin-5-ylidenemethyl)benzoic acid amide and the like; the compounds described in WO99/15129, preferably 4-{4-[2-(2-[bis(4-chlorophenyl)methoxy]ethylsulfonyl)ethoxy]phenyl}-1,1,1-trifluoro-2-butanone and the like; the compounds described in WO98/05637, preferably 1-{2-[4-(carboxymethyl)phenoxy]ethyl}-3-dodecanoylindole-2-carboxylic acid and the like; the compounds described in Japanese Patent Unexamined Publication (Kokai) No. 2002-80368, preferably 4-methyl-2-oxo-5-(5,6,7,8-tetrahydronaphthalen-2-yl)oxazolidine-3-carboxylic acid (6-methoxytetrahydropyran-2-yl)amide, 4-methyl-2-oxo-5-(4-methylphenyl)thiazolidine-3-carboxylic acid (tetrahydropyran-2-yl)amide

and the like; and the type 4 PLA<sub>2</sub> inhibitors selected from the compounds described in WO98/08818, the compounds described in WO99/43651, the compounds described in WO99/43672, the compounds described in WO03/048122, the compounds described in WO95/10508, the compounds described in WO97/05135, the compounds described in Japanese Patent Unexamined Publication No. 7-126166, the compounds described in Japanese Patent Unexamined Publication No. 7-224076, the compounds described in Japanese Patent Unexamined Publication No. 7-224076, the compounds described in Japanese Patent Unexamined Publication No. 2000-119292, the compounds described in Japanese Patent Unexamined Publication No. 2000-109432, the compounds described in Japanese Patent Unexamined Publication No. 7-223997, the compounds described in the U.S. Patent No. 5,994,398, the compounds described in WO00/27824, the compounds described in Japanese Patent Unexamined Publication No. 2000-38380, the compounds described in WO00/71118, the compounds described in Japanese Patent No. 3107613, the compounds described in WO03/031414, the compounds described in U.S. Patent No. 5,453,443, and the compounds described in WO02/038575. Examples further include the following known type 4 PLA<sub>2</sub> inhibitors described in references: arachidonyl trifluoromethyl ketone (Street et al., *Biochemistry*, 1993, vol. 32, p.5935); methyl arachidonyl fluorophosphate (Kennedy et al., *Mediators of Inflammation*, 1994, vol. 3, p.337);  $\beta$ -lactam derivatives (Burke et al., *J. Enzyme Inhibition*, 1998, vol. 13, p.195); choline derivatives (Burke et al., *J. Biol.Chem.*, 1999, vol. 274, p.18864); 1,3-disubstituted propan-2-one derivatives, especially 4-[3-(4-decyloxyphenyloxy)-2-oxopropoxy]benzoic acid (Connolly et al., *J. Med.Chem.*, 2002, vol. 45, p.1348); Surfactin (Kim et al., *Biochem. Pharmacol.*, 1998, vol. 55, p.975); 1,1,1-trifluorononadeca-10,13,16-trien-2-one and 1,1,1-trifluorononadeca-10,13-dien-2-one (Amandi-Burgermeister et al., *Eur. J. Pharmacol.*, 1997, vol. 326, p.237); and 2-oxoamide derivatives (Kokotos et al., *J. Med. Chem.*, 2002, vol. 45, p.2891).

In the present invention, preferred examples of type 4 PLA<sub>2</sub> inhibitor further include the compounds represented by the aforementioned formula (I) and pharmacologically acceptable salts thereof. Various combinations of the compounds represented by the formula (I) and pharmacologically acceptable salts thereof described in the specification can also be arbitrarily chosen.

When a medicament comprising a type 4 PLA<sub>2</sub> inhibitor as an active ingredient is used as a prophylactic and/or therapeutic agent for fibrosis, as for Compound (I) of the present invention, for example, Compound (I) of the present invention or a pharmaceutically acceptable salt thereof, per se, may be used in an effective amount, or the substance may be used after preparation of a pharmaceutical composition in the form of solid, liquid or gel by mixing the substance with a pharmaceutically acceptable carrier. As for the pharmaceutically acceptable carrier, known information and the information about carriers described in this specification can be referred to. As for known type 4 PLA<sub>2</sub> inhibitors, a known type 4 PLA<sub>2</sub> inhibitor or a pharmaceutically acceptable salt thereof, per se, may be used in an effective amount, or as mentioned above, the inhibitors may be used after preparation of a pharmaceutical composition by mixing the inhibitor with a pharmaceutically acceptable carrier.

It would be readily understood by those skilled in the art that progression-preventing agents, that is used for preventing progression of pathological conditions, occasionally fall within the scope of the agent for prophylactic and/or therapeutic treatment of the present invention.

Examples of the dosage form for preparation of the aforementioned pharmaceutical composition, tablet, powder, granule, syrup, suspension, capsule, inhalant, injection, and the like, and in order to prepare the compositions, various carriers are used depending on the type of the composition. Examples of the carrier for oral agents include, for example, excipients, binders, lubricants,

flowability improvers, and colorants. When an inhalant is prepared (examples of administration method include a method of inhaling powder of the pharmaceutical composition or a solution obtained by dissolving or suspending the pharmaceutical composition in a solvent, per se, a method of inhaling mist of the composition prepared by using a sprayer called atomizer or nebulizer), the preparation the aforementioned pharmaceutical composition in the form of solid can be referred to for preparation of a powder for the inhalation, and a powder obtained is preferably further made into micropowder. When the composition is inhaled as a liquid, preferred examples of the preparation method include a method of dissolving a solid pharmaceutical composition, which is prepared by referring to the above explanation, in distilled water or a suitable solvent to obtain a solution of medicament upon use, and a method of preparing a liquid pharmaceutical composition prepared by referring the above explanation to obtain a solution of medicament. As for a size of the aforementioned powder or mist of a solution of a medicament to be inhaled, a particle size may be suitable for inhalation. For example, an upper limit is preferably  $100\ \mu\text{m}$  or less, further preferably  $50\ \mu\text{m}$  or less, most preferably  $10\ \mu\text{m}$  or less. A lower limit is not particularly limited, and a smaller particle size is more preferred. When an injection and the like are prepared, distilled water for injection, physiological saline, glucose solution, vegetable oil for injection, propylene glycol, polyethylene glycols and the like can generally be used as diluents. Further, antimicrobial agents, antiseptics, stabilizers, isotonic agents, soothing agents, and the like may be added, as required.

When the aforementioned prophylactic and/or therapeutic agent is administered, a suitable dosage form can be chosen and administered via a suitable route. For example, the agent can be orally administered in the form of a tablet, a powder, a granule, a syrup, a suspension, or a capsule. The agent can also be administered via transairway route in the form of an inhalant. Further, the agent

can be administered subcutaneously, intradermally, intravascularly, intramuscularly or intraperitoneally in the form of injection including a drip infusion. Furthermore, the agent can be transmucosally administered in the form of a sublingual agent or a suppository, and can be transdermally administered in the form of a gel, a lotion, an ointment, a cream, or a spray.

A dose thereof varies depending on the dosage form, and the age, weight, degree of symptoms of a patient and the like. Examples of the dose include generally an administration at a dose of 1 to 1,000 mg per day for an adult once to three times a day. Every day administration for a period of several days to two months is commonly applied. The daily dose and the administration period may be increased or decreased depending on symptoms of a patient.

As for the application of the aforementioned prophylactic and/or therapeutic agent, the agent may be administered to patients with pulmonary fibrosis as explained above. In addition, the prophylactic and/or therapeutic agent of the present invention containing a type PLA<sub>2</sub> inhibitor as an active ingredient may preferably be administered after the administration of, most preferably immediately after the administration of an agent, which may possibly induces pulmonary fibrosis as an adverse reaction. Furthermore, as for the administration time, the prophylactic and/or therapeutic agent of the present invention may be administered simultaneously with an agent which may possibly induces pulmonary fibrosis as an adverse reaction, or the agent of the present invention may be administered beforehand.

## Examples

The present invention will be further specifically explained with reference to examples. However, the scope of the present invention is not limited to the following examples. In the examples, for thin layer chromatography (TLC),

Precoated Silica Gel 60 F254 (produced by Merck, product number: 5715-1M)) was used. After development with chloroform:methanol (1:0 to 1:1), acetonitrile:acetic acid:water (200:1:1 to 100:4:4) or ethyl acetate:hexane (1:0 to 0:1), spots were observed by UV irradiation (254 nm) or color development with ninhydrine or dinitrophenylhydrazine solution in hydrochloric acid. For drying organic solvent, anhydrous magnesium sulfate or anhydrous sodium sulfate was used. As for column chromatography, the indication of "Quad" means use of Quad 1 preparative chromatography system (produced by Biotage), and one or several columns selected from cartridge columns KP-Sil-12M, 40S and 40M produced by the same manufacturer were used depending on the amount of sample. For flash column chromatography, Silica gel 60N (spherical shape, neutral, 40 to 100  $\mu$  m, produced by Kanto Chemicals) was used. Preparative thin layer chromatography (hereinafter abbreviated as "PTLC") was performed by using one or several plates of PLC Plate Silica Gel 60 F254 (20 x 20 cm, thickness: 2 mm, concentration zone: 4 cm, produced by Merck, product number: 13793-1M) were used depending on the amount of sample.

The indication of "LCMS" means that mass spectrum was measured by liquid chromatography-mass spectrometry (LC-MS). Platform-LC type mass spectrometry apparatus (produced by Micromass) was used as the mass spectrometer, and the measurement was performed by the electrospray ionization (ESI) method. As a liquid chromatography apparatus, an apparatus produced by GILSON was used. As a separation column, Mightysil RP-18 GP 50-4.6 (produced by Kanto Chemicals) was used. Elution was generally performed at a flow rate of 2 ml/minute, and Solution A = water [containing 0.1% (v/v) acetic acid] and Solution B = acetonitrile [containing 0.1% (v/v) acetic acid] were used as solvents.

In the tables mentioned below, data indicated by "LCMS" mean data of liquid chromatography-mass spectrometry spectra. In the columns of "Mass", data

of mass spectrometry were shown (the indication "N.D" means that no molecular ion peak was detected). In the columns of "method", elution conditions of the liquid chromatography are described. In the columns of "RTime", retention times in the liquid chromatography are shown. For the indication of retention time in the liquid chromatography, the indication "A" for elution condition means that measurement was performed by elution with a linear gradient of 5 to 100% (v/v) Solution B from 0 minute to 5 minutes and then with 100% Solution B until 6 minutes. Similarly, the indication "B" for elution condition means that measurement was performed by elution with 30% (v/v) Solution B from 0 minute to 0.5 minute, then with a linear gradient of 30 to 95% (v/v) Solution B from 0.5 minute to 4 minutes and then with 95% (v/v) Solution B until 6 minutes. For the compounds with the indication C in the columns of elution conditions, data of mass spectrometry measured by fast atomic bombardment mass spectrometry (FAB-MS) using JEOL-JMS-SX102 (produced by JEOL Co., Ltd.) were mentioned in the columns of "Mass". Further, for the compounds with the indication D in the elution conditions, an apparatus manufactured by Waters Ltd. was used as a liquid chromatography apparatus. As a column for separation, Develosil C30-UG-5 (50 x 4.6 mm, Nomura Kagaku Co., Ltd.) was used. Measurement was performed under elution condition with a linear gradient of 5 to 98% (v/v) Solution B from 0 minute to 4 minutes and then with 100% Solution B until 6 minutes.

In the columns indicated as "Exp.", compound numbers are shown. When the tables include a column indicated as "position", substituting positions of substituents are indicated in the column. The abbreviations used in the tables have the following meanings.

n: normal, i: iso, s: secondary, t: tertiary, c: cyclo, D: di, Me: methyl, Et: ethyl, Pr: propyl, Bu: butyl, Pen: pentyl, Hex: hexyl, Hep: heptyl, Ph: phenyl, Bn: benzyl, Py: pyridyl, Indan: indanyl, Ac: acetyl, CHO: formyl, COOH: carboxyl, NO<sub>2</sub>: nitro,

DMA: dimethylamino, NH<sub>2</sub>: amino, CF<sub>3</sub>: trifluoromethyl, F: fluoro, Cl: chloro, Br: bromo, OMe: methoxy, OH: hydroxy, TFA: trifluoroacetyl, SO<sub>2</sub>: sulfonyl, CO: carbonyl, Nap: naphthyl, Ind: 1H-indolyl, 1HIdz: 1H-indazolyl, 2HIdz: 2H-indazolyl, Bzt: benzothiazole, 2ABzt: 2-aminobenzothiazole, BF: benzofuranyl, BT: benzo[b]thienyl, Qu: Quinolyl, IQ: isoquinolyl

The numbers given before the substituents indicate substituting positions. The numbers given with hyphens before abbreviations of aromatic rings indicate substituting positions of the aromatic rings. (S) indicates optically active substances with S-configuration, and (R) indicates optically active substances with R-configuration. Representative examples of the substituents shown in the tables with abbreviations are listed in Table 2 mentioned below.

Table 2

Structure	abbreviation	Structure	abbreviation	Structure	abbreviation
	cPenMeO		cHexMeO		iBuO
	2EtBuO		2,3DMeBuO		cPenO
	cHexO		cHepO		BnO
	(R)1PhEtO		2ClBnO		4FBnO
	2-IndanO		2(4FPh)EtO		2(4DMAPh)EtO
	2(3-Py)EtO		2(PhO)EtO		3F,4(OMe)BnO
			2-Nap		1-Nap
	5-Ind		1Me-5-Ind		5-1HIdz
	1Me-5-1HIdz		5-Bzt		5-2ABzt
	2Me-5-Bzt		5-BT		5-BF
	3-Qu		6-IQ		



The manufacturers of the reagents used may sometimes be indicated with the following abbreviations.

TCI: Tokyo Kasei Kogyo Co., Ltd., Ald: Aldrich Co., KANTO: Kanto Kagaku, WAKO: Wako Pure Chemical Industries, Ltd., LANC: Lancaster Synthesis, MAYB: Maybridge, plc.

[Example A-1]

Synthesis of methyl 3-(4-hydroxyphenyl)propionate (Intermediate 1)

A solution obtained beforehand by adding thionyl chloride (18.3 ml, WAKO) dropwise to methanol (250 ml) and mixing the mixture under ice cooling was added dropwise with a solution of 3-(4-hydroxyphenyl)propionic acid (16.6g, TCI) in methanol (50 ml) under ice cooling, stirred for 30 minutes, warmed to room temperature, and further stirred for 1.5 hours. The reaction mixture was concentrated under reduced pressure, and then extracted with diethyl ether (200 ml). The organic layer was washed successively with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride and saturated brine. The organic layer was dried, and then the solvent was evaporated under reduced pressure to obtain the title compound (Intermediate 1, 17.95 g).

Synthesis of methyl 3-(4-cyclopentylmethoxyphenyl)propionate (Intermediate 2)

A solution of cyclopentane methanol (4.05 ml, Ald) in anhydrous tetrahydrofuran (abbreviated as "THF" hereinafter, 40 ml) was added with triethylamine (6.49 ml, WAKO), added dropwise with methanesulfonyl chloride (3.48 ml, WAKO) under ice cooling, and stirred for 30 minutes. The reaction mixture was added with water (50 ml), and extracted with diethyl ether (80 ml x 2). The organic layer was washed with saturated brine and dried, and then the solvent was evaporated under reduced pressure. A solution obtained beforehand by adding 60% sodium hydride (1.15 g, KANTO) to a solution of Intermediate 1 (4.50 g) in

N,N-dimethylformamide (abbreviated as "DMF" hereinafter, 35 ml) under ice cooling and stirring the solution for 15 minutes was added with a solution of the aforementioned residue in DMF (10 ml) under ice cooling. The reaction mixture was stirred for 15 minutes, then warmed to room temperature, stirred for 45 minutes, and further stirred at 60°C for 15 hours. The reaction mixture was added with water (100 ml) and diethyl ether (200 ml) for extraction. The organic layer was successively washed with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride, and saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography (hexane:isopropyl ether = 9:1) to obtain the title compound (Intermediate 2, 5.58 g).

Synthesis of methyl 3-(3-bromo-4-cyclopentylmethoxyphenyl)propionate  
(Compound No. A-1)

A solution of Intermediate 2 (1.31 g) in acetonitrile (50 ml) was added with N-bromosuccinimide (hereinafter abbreviated as "NBS", 979 mg, KANTO), stirred at room temperature for 2 hours, then warmed to 40°C, and stirred for 3 hours. The reaction mixture was concentrated under reduced pressure, then added with ethyl acetate (200 ml) and washed successively with saturated aqueous ammonium chloride, 5% aqueous sodium sulfite, saturated aqueous sodium hydrogencarbonate and saturated brine. The organic layer was dried, and then the solvent was evaporated under reduced pressure to obtain the title compound (Compound No. A-1, 1.69 g).

[Example A-2]

Synthesis of 3-(3-bromo-4-methoxyphenyl)propionic acid (Intermediate 3)

According to the procedure described in the synthesis method of Compound No. A-1 provided that the reaction was carried out under ice cooling for 30 minutes and at room temperature for 3 hours, 3-(4-methoxyphenyl)propionic acid (27.0 g,

TCI) and NBS (29.4 g) were reacted and treated to obtain the title compound (Intermediate 3, 38.1 g).

#### Synthesis of 3-(3-bromo-4-hydroxyphenyl)propionic acid (Intermediate 4)

According to a procedure described in a literature (Carreno, M.C., J. Org. Chem., 1995, vol. 60, p.5328), a 1 M solution of boron tribromide in methylene chloride (200 ml, Fluka) was added dropwise with a solution of Intermediate 4 (23.5 g) in methylene chloride (200 ml) at -78°C, warmed to room temperature after 30 minutes, and further stirred for 1.5 hours. The reaction mixture was poured into ice water (750 ml), and stirred at room temperature for 1 hour. The reaction mixture was added with diethyl ether (750 ml) for extraction. The organic layer was added with 2 N aqueous sodium hydroxide (250 ml x 2) for extraction, and then the aqueous layer was made acidic with 5 N aqueous hydrochloric acid under ice cooling, and extracted with diethyl ether (375 ml x 2) again. The organic layer was washed with saturated brine and dried, and then the solvent was evaporated under reduced pressure. The organic layer was washed with saturated brine and dried, and then the solvent was evaporated under reduced pressure to obtain the title compound (Intermediate 4, 23.5 g).

#### Synthesis of methyl 3-(3-bromo-4-hydroxyphenyl)propionate (Intermediate 5)

According to the procedure described in the synthesis method of Intermediate 1 provided that the purification was performed by flash column chromatography (hexane:ethyl acetate = 4:1), Intermediate 4 (21.15 g) and thionyl chloride (15.0 ml) were reacted and treated in methanol to obtain the title compound (Intermediate 5, 20.36 g).

#### Synthesis of methyl (3-bromo-4-cyclohexylmethoxyphenyl)propionate (Compound No. A-2)

A solution of Intermediate 5 (1.29 g) in DMF (25 ml) was added with potassium carbonate (0.86 g) and bromomethylcyclohexane (1.05 ml, TCI), stirred

under argon atmosphere at room temperature for 2 hours, then warmed to 60°C, and stirred for 17 hours. The reaction mixture was poured into ice water, and extracted with isopropyl ether (200 ml). The organic layer was successively washed with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride, and saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography (hexane:isopropyl ether = 9:1) to obtain the title compound (Compound No. A-2, 1.45 g).

[Example A-5]

Synthesis of methyl 3-(3-bromo-4-cyclopentyloxyphenyl)propionate (Compound No. A-5)

A solution of Intermediate 5 (4.50 g) in DMF (20 ml) was added with 60% sodium hydride (440 mg, KANTO) under ice cooling. The reaction mixture was stirred for 10 minutes, then added with bromocyclopentane (1.61 ml, TCI), warmed to room temperature, stirred for 1.5 hours, then warmed to 60°C, and further stirred for 16 hours. The reaction mixture was added with water (50 ml) and isopropyl ether (300 ml) for extraction. The organic layer was successively washed with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride, and saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography (hexane:isopropyl ether = 7:1) to obtain the title compound (Compound No. A-5, 2.50 g).

[Example A-6]

Synthesis of methyl 3-(3-bromo-4-cyclohexyloxyphenyl)propionate (Compound No. A-6)

A solution of Intermediate 5 (2.06 g), triphenylphosphine (hereinafter abbreviated as "Ph<sub>3</sub>P", 6.28 g, WAKO) and cyclohexanol (2.53 ml, WAKO) in

anhydrous THF (60 ml) was added dropwise with a 40% solution of diisopropylazodicarboxylic acid ester in toluene (hereinafter abbreviated as "40% DIAD", 11.35 ml, WAKO) under ice cooling over 10 minutes. The reaction mixture was stirred for 10 minutes, then warmed to room temperature, and stirred for 18.5 hours. The reaction mixture was added with water (50 ml) and ethyl acetate (200 ml)) for extraction. The organic layer was successively washed with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride and saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:isopropyl ether = 8:1) to obtain the title compound (Compound No. A-6, 2.35 g).

[Example A-20]

Synthesis of methyl 3-(3-bromo-5-chloro-4-hydroxyphenyl)propionate (Intermediate 6)

A solution of Intermediate 5 (516mg) in chloroform (5 ml) was added with sulfuryl chloride (177  $\mu$ l), and stirred at room temperature for 21 hours. The reaction mixture was poured into aqueous saturated sodium hydrogencarbonate (20 ml), and extracted with ethyl acetate. The organic layer was successively washed with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride, and saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 10:1) to obtain the title compound (Intermediate 6, 290mg).  
Synthesis of methyl 3-(3-bromo-5-chloro-4-cyclopentylmethoxyphenyl)propionate (Compound No. A-20)

According to the procedure described in the synthesis method of Compound No. A-6 provided that the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 30:1), Intermediate 6 (278 mg), Ph<sub>3</sub>P (747 mg),

cyclopentane methanol (308  $\mu$  l), and 40% DIAD (1.34 ml) were reacted and treated to obtain the title compound (Compound No. A-20, 337mg).

[Example A-21]

Synthesis of ethyl 3-(3-fluoro-4-methoxyphenyl)acrylate (Intermediate 7)

A solution of 3-fluoro-4-methoxybenzaldehyde (2.20 g, Ald) in 1,2-diethoxyethane (5 ml) was added with ethyl diethylphosphonoacetate (3.12 ml, TCI) and added with 60% sodium hydride (624mg) under ice cooling. After being stirred for 10 minutes, the reaction mixture was warmed to room temperature, and stirred for 5 hours. The reaction mixture was added with ethyl acetate (90 ml), and washed successively with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride and saturated brine. The organic layer was dried, and then the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography (Quad, hexane:ethyl acetate = 10:1) to obtain the title compound (Intermediate 7, 3.16 g).

Synthesis of ethyl 3-(3-fluoro-4-methoxyphenyl)propionate (Intermediate 8)

A solution of Intermediate 7 (3.01 g) in ethyl acetate (50 ml) and methanol (25 ml) was added with 10% palladium/carbon (300 mg, Merck), and stirred at room temperature for 2 hours under hydrogen atmosphere. The reaction mixture was filtered, and the solvent of the filtrate was evaporated under reduced pressure to obtain the title compound (Intermediate 8, 3.02 g).

Synthesis of 3-(3-fluoro-4-methoxyphenyl)propionic acid (Intermediate 9)

A solution of Intermediate 8 (2.97 g) in methanol (40.0 ml) was added with 2 N aqueous sodium hydroxide (15.0 ml) and stirred at 60°C for 16 hours. The reaction mixture was concentrated under reduced pressure, then made acidic with aqueous 5% hydrochloric acid under ice cooling, and extracted with ethyl acetate (200 ml). The organic layer was washed with saturated brine and dried, and then the solvent was evaporated under reduced pressure to obtain the title compound

(Intermediate 9, 2.40 g).

#### Synthesis of 3-(3-fluoro-4-hydroxyphenyl)propionic acid (Intermediate 10)

A pyridine/hydrochloric acid complex prepared by mixing pyridine (30 ml) and concentrated hydrochloric acid (30 ml) and heating the mixture at 190°C for 1 hour was added with Intermediate 9 (2.40 g) and stirred at 190°C for 1.5 hours. The reaction mixture was poured into 1 N hydrochloric acid (100 ml) cooled with ice, and extracted with ethyl acetate (200 ml). The organic layer was washed with saturated brine and dried, and then the solvent was evaporated under reduced pressure to obtain the title compound (Intermediate 10, 1.98 g).

#### Synthesis of methyl 3-(3-fluoro-4-hydroxyphenyl)propionate (Intermediate 11)

According to the procedure described in the synthesis method of Intermediate 1, Intermediate 10 (1.77 g) and thionyl chloride (1.65 ml) were reacted and treated in methanol to obtain the title compound (Intermediate 11, 1.85 g).

#### Synthesis of methyl 3-(3-bromo-5-fluoro-4-hydroxyphenyl)propionate (Intermediate 12)

According to the procedure described in the synthesis method of Compound No. A-1 with the modifications that the reaction was carried out for 2 hours under ice cooling, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 10:1), Intermediate 11 (1.84 g) and NBS (1.74 g) were reacted and treated to obtain the title compound (Intermediate 12, 1.74 g).

#### Synthesis of methyl 3-(3-bromo-4-cyclopentylmethoxy-5-fluorophenyl)propionate (Compound No. A-21)

According to the procedure described in the synthesis method of Compound No. A-6 with the modifications that the reaction was carried out for 22 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 50:1), Intermediate 11 (310 mg), tributylphosphine (hereinafter abbreviated as "tBu<sub>3</sub>P", 405  $\mu$ l, WAKO) instead of Ph<sub>3</sub>P, cyclopentane methanol

(176  $\mu$  l), and N,N,N',N'-tetramethylazodicarboxamide (hereinafter abbreviated as "TMAD", 279 mg, TCI) instead of 40% DIAD were reacted and treated to obtain the title compound (Compound No. A-21, 386 mg).

[Example A-24]

Synthesis of 4-cyclopentyloxy-3-methylbenzaldehyde (Intermediate 13)

According to the procedure described in the synthesis method of Compound No. A-2 with the modifications that the reaction was carried out for 16 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 9:1), 4-hydroxy-3-methylbenzaldehyde (283 mg, TCI), potassium carbonate (578 mg) and bromocyclopentane (430  $\mu$  l) were reacted and treated to obtain the title compound (Intermediate 13, 350 mg).

Synthesis of ethyl 3-(4-cyclopentyl-3-methylphenyl)acrylate (Intermediate 14)

According to the procedure described in the synthesis method of Intermediate 7 with the modifications that the reaction was carried out for 2 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 9:1), Intermediate 13 (342 mg), ethyl diethylphosphonoacetate (408  $\mu$  l) and 60% sodium hydride (82 mg) were reacted and treated to obtain the title compound (Intermediate 14, 450 mg).

Synthesis of ethyl 3-(4-cyclopentyl-3-methylphenyl)propionate (Intermediate 15)

According to the procedure described in the synthesis method of Intermediate 8, Intermediate 14 (446 mg) and 10% palladium/carbon (20 mg) were reacted and treated under hydrogen gas atmosphere to obtain the title compound (Intermediate 15, 439 mg).

Synthesis of ethyl 3-(3-bromo-4-cyclopentyl-5-methylphenyl)propionate (Compound No. A-24)

According to the procedure described in the synthesis method of Compound No. A-1, Intermediate 15 (437 mg) and NBS (320 mg) were reacted and treated to



obtain the title compound (Compound No. A-24, 545 mg).

[Example A-25]

Synthesis of 3-bromo-4-(t-butyldimethylsilyloxy)-5-methoxybenzaldehyde  
(Intermediate 16)

A solution of 3-bromovanillin (1.16 g, TCI) in anhydrous DMF (20 ml) was added with imidazole (408 mg, TCI), added dropwise with a solution of 4-(N,N-dimethylamino)pyridine (25 mg) and t-butyldimethylsilyl chloride (904 mg, TCI) in DMF (15 ml) under ice cooling, stirred 30 minutes, then warmed to room temperature, and further stirred 3 hours. The reaction mixture was added with water (100 ml), and extracted with ethyl acetate (100 ml). The organic layer was washed with saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography (hexane:ethyl acetate = 9:1) to obtain the title compound (Intermediate 16, 1.75 g).

Synthesis of ethyl 3-[3-bromo-4-(t-butyldimethylsilyloxy)-5-methoxyphenyl]acrylate  
(Intermediate 17)

According to the procedure described in the synthesis method of Intermediate 7 with the modifications that the reaction was carried out for 1.5 hours, and the purification was performed by flash column chromatography (hexane:ethyl acetate = 9:1), Intermediate 16 (910 mg), ethyl diethylphosphonoacetate (530  $\mu$ l) and 60% sodium hydride (120 mg) were reacted and treated to obtain the title compound (Intermediate 17, 937 mg).

Synthesis of ethyl 3-[3-bromo-4-(t-butyldimethylsilyloxy)-5-methoxyphenyl]propionate (Intermediate 18)

According to the procedure described in the synthesis method of Intermediate 8, Intermediate 17 (945 mg) and 10% palladium/carbon (95 mg) were reacted and treated under hydrogen gas atmosphere to obtain the title compound (Intermediate 18, 760 mg).

Synthesis of ethyl 3-(3-bromo-4-hydroxy-5-methoxyphenyl)propionate (Intermediate 19)

A solution of Intermediate 18 (750 mg) in THF (50 ml) was added with a 1 M solution of tetrabutylammonium fluoride in THF (5 ml, TCI), and stirred for 1.5 hours. The reaction mixture was added with saturated aqueous sodium hydrogencarbonate (30 ml), and extracted with ethyl acetate (50 ml). The organic layer was washed with saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography (hexane:ethyl acetate = 4:1) to obtain the title compound (Intermediate 19, 542 mg).

Synthesis of ethyl 3-(3-bromo-4-cyclopentyloxy-5-methoxyphenyl)propionate (Compound No. A-25)

According to the procedure described in the synthesis method of Compound No. A-6 with the modifications that the reaction was carried out for 16 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 7:1), Intermediate 19 (400 mg),  $\text{Ph}_3\text{P}$  (1.31 g), cyclopentanol (450  $\mu\text{l}$ ), and TMAD (860 mg) were reacted and treated to obtain the title compound (Compound No. A-25, 376 mg).

[Example A-26]

Synthesis of methyl 3-(3-bromo-4-cyclopentylmethoxy-5-nitrophenyl)propionate (Compound No. A-26)

A solution obtained beforehand by adding 70% nitric acid (3.9 ml) to acetic anhydride (30 ml) under ice cooling and stirring the mixture for 10 minutes was added with a solution of Compound No. A-1 (5.12 g) in acetonitrile (25 ml) at  $-15^\circ\text{C}$  over 15 minutes, and stirred further for 15 minutes. The reaction mixture was poured into 1 N aqueous sodium hydroxide (500 ml) containing ice, and extracted with diethyl ether (300 ml x 2). The organic layer was successively washed with

saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride, and saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 10:1) to obtain the title compound (Compound No. A-26, 3.68 g).

[Example A-31]

Synthesis of methyl 3-(3-bromo-4-phenoxyphenyl)propionate (Compound No. A-31)

A solution of Intermediate 5 (3.08 g) in anhydrous N-methylpyrrolidone (9.5 ml, WAKO) was successively added with cesium carbonate (3.58 g, WAKO), iodobenzene (1.4 ml, TCI), dipivaloylmethane (0.12 ml, TCI) and copper(I) chloride (275 mg, WAKO), and stirred 120°C for 16 hours under argon gas atmosphere. The reaction mixture was added with t-butyl methyl ether (25 ml), and insoluble solids were removed by filtration. The filtrate was washed successively with 2 N aqueous hydrochloric acid and saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography (hexane:ethyl acetate = 1:10) to obtain the title compound (Compound No. A-31, 1.00 g).

[Example B-96]

Synthesis of methyl 3-(3-bromo-4-methoxyphenyl)propionate (Intermediate 20)

According to the procedure described in the synthesis method of Intermediate 1 provided that the purification was performed by flash column chromatography (hexane:ethyl acetate = 6:1), Intermediate 3 (1.60 g) and thionyl chloride (1.44 ml) were reacted and treated in methanol to obtain the title compound (Intermediate 20, 1.63 g).

Synthesis of methyl 3-(3-bromo-4-methoxy-5-nitrophenyl)propionate (Intermediate 21)

A solution of Intermediate 20 (3.20 g) in acetic anhydride (25 ml) was added

with potassium nitrate (1.30 g) under ice cooling and stirred for 10 minutes, and the solution was added dropwise with concentrated sulfuric acid (730  $\mu$ l) over 10 minutes. The reaction mixture was stirred for 10 minutes for 10 minutes at the same temperature, then warmed to room temperature, and further stirred for 30 minutes. The reaction mixture was poured into 1 N aqueous sodium hydroxide (250 ml) containing ice, and extracted with isopropyl ether (200 ml x 2). The organic layer was successively washed with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride, and saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 10:1) to obtain the title compound (Intermediate 21, 2.73 g).

Synthesis of 3-(3-bromo-4-methoxy-5-nitrophenyl)propionic acid (Intermediate 22)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 1 hour, Intermediate 21 (12.73 g) and 2 N aqueous sodium hydroxide (40 ml) were reacted and treated to obtain the title compound (Intermediate 22, 11.53 g).

Synthesis of 3-(3-bromo-4-hydroxy-5-nitrophenyl)propionic acid (Intermediate 23)

According to the procedure described in the synthesis method of Intermediate 4 provided that the reaction was carried out for 2 hours, Intermediate 22 (11.53 g) and a 1 M solution of boron tribromide in methylene chloride (100 ml) were reacted and treated to obtain the title compound (Intermediate 23, 10.68 g).

Synthesis of methyl 3-(3-bromo-4-hydroxy-5-nitrophenyl)propionate (Intermediate 24)

According to the procedure described in the synthesis method of Intermediate 1 provided that the reaction was carried out for 17.5 hours, Intermediate 23 (10.68 g) and thionyl chloride (8.06 ml) were reacted and treated to obtain the title compound (Intermediate 24, 8.27 g).

Synthesis of methyl 3-[3-bromo-4-(indan-2-yloxy)-5-nitrophenyl]propionate  
(Compound No. B-96)

According to the procedure described in the synthesis method of Compound No. A-6 with the modifications that the reaction was carried out for 15 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 19:1), Intermediate 24 (151 mg), Ph<sub>3</sub>P (260 mg), 2-hydroxyindane (133 mg, TCI) and 40% DIAD (470  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. B-96, 192 mg).

[Example B-99]

Synthesis of methyl 3-(3-amino-5-bromo-4-cyclopentyloxyphenyl)propionate  
(Compound No. B-99)

A solution of Compound No. A-28 (416 mg) in a mixture of THF (5 ml) and methanol (5 ml) was added with Raney 2800 nickel (230 mg, Ald) and stirred at room temperature for 6 hours under hydrogen atmosphere. The reaction mixture was filtered, and the solvent of the filtrate was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 5:2) to obtain the title compound (Compound No. B-99, 143 mg).

[Example B-103]

Synthesis of methyl 3-[4-benzyloxy-5-bromo-3-(2,2,2-trifluoroacetyl-amino)phenyl]propionate (Compound No. B-103)

A solution of Compound No. B-100 (58.7 mg) in methylene chloride (2 ml) was added with triethylamine (76  $\mu$ l), added dropwise trifluoroacetic anhydride (91  $\mu$ l, TCI) under ice cooling, stirred for 30 minutes, then warmed to room temperature, and further stirred for 2 hours. The reaction mixture was added with water (5 ml), and extracted with methylene chloride (20 ml). The organic layer was washed with saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column

chromatography (Quad, hexane:ethyl acetate = 3:1) to obtain the title compound (Compound No. B-103, 59.1 mg).

[Example B-105]

Synthesis of methyl 3-[4-benzyloxy-5-bromo-3-(N-methylamino)phenyl]propionate (Compound No. B-105)

A solution of Compound No. B-100 (105 mg) in DMF (3 ml) was added with 60% sodium hydride (20 mg) under ice cooling, and stirred for 10 minutes. This reaction mixture was added dropwise with methyl iodide (32  $\mu$ l), stirred for 10 minutes, then warmed to room temperature, and further stirred for 2 hours. The reaction mixture was poured into water, and added with ethyl acetate (30 ml) for extraction. The organic layer was successively washed with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride, and saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 6:1) to obtain the title compound (Compound No. B-105, 17 mg).

[Example B-109]

Synthesis of 3-[4-benzyloxy-5-bromo-3-(N,N-dimethylamino)phenyl]propionic acid (Compound No. B-109)

A solution of Compound No. B-100 (105 mg) in DMF (3 ml) was added with 60% sodium hydride (40 mg) under ice cooling, and stirred for 10 minutes. This reaction mixture was added dropwise with methyl iodide (300  $\mu$ l), stirred for 10 minutes, then warmed to room temperature, and further stirred for 16 hours. The reaction mixture was poured into water, and added with ethyl acetate (30 ml) for extraction. The organic layer was successively washed with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride, and saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate =

6:1) to obtain the title compound (Compound No. B-109, 88 mg).

[Examples B-113 and B-114]

Syntheses of 3-(3-bromo-4-cyclopentyloxy-5-hydroxyphenyl)propionic acid (Compound No. B-113) and 3-(5-acetoxy-3-bromo-4-cyclopentyloxyphenyl)propionic acid (Compound No. B-114)

A solution of Compound No. B-99 (415 mg) in acetic acid (1.5 ml) was added with 20% sulfuric acid (1.0 ml). This reaction mixture was added dropwise with an aqueous solution (0.5 ml) of sodium nitrite (78 mg) over 10 minutes, while the temperature of the reaction mixture was maintained below 10°C, and further stirred for 5 minutes. This reaction mixture was added dropwise to a solution of sodium acetate (348 mg) in acetic acid (3.5 ml) heated and stirred at 100°C beforehand over 5 minutes, and further stirred for 10 minutes with heating. The reaction solution was poured into ice water (50 ml), and extracted with isopropyl ether (100 ml x 2). The organic layer was successively washed with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride and saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 10:1) to obtain the title compounds (Compound No. B-113, 47mg and Compound No. B-114, 105 mg).

[Example B-117]

Synthesis of methyl 3-(3,5-dibromo-4-cyclopentylmethoxyphenyl)propionate (Compound No. B-117)

A solution of Intermediate 1 (670 mg) in acetonitrile (30 ml) was added with NBS (990 mg), stirred at room temperature for 2 hours, then warmed to 40°C, and stirred for 18 hours. The reaction mixture was concentrated under reduced pressure, then added with ethyl acetate (100 ml), and washed successively with saturated aqueous ammonium chloride, 5% aqueous sodium sulfite, saturated

aqueous sodium hydrogencarbonate and saturated brine. The organic layer was dried, and then the solvent was evaporated under reduced pressure. According to the procedure described in the synthesis method of Compound No. A-6 with the modifications that the reaction was carried out for 18 hours under ice cooling, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 10:1), the residue was reacted with  $\text{Ph}_3\text{P}$  (1460 mg), cyclopentane methanol (560 mg) and 40% DIAD (2.6 ml) and treated to obtain the title compound (Compound No. B-117, 710 mg).

[Examples A-1 to A-33]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification including the examples described above are shown in Table-A-1. The compounds were prepared according to the preparation methods of the compound numbers (e.g., "A-1") or the intermediate numbers (e.g., "Int 2") shown in the columns of "Syn" in the tables. "Int" means an intermediate compound number. When the preparation required a plurality of steps, a plurality of compound numbers or intermediate compound numbers are mentioned in the columns of "Syn". For example, an indication of "Int 2, A-1" in a column of "Syn" means that "the compound is prepared from a compound prepared according to the procedure described in the synthesis method of Intermediate 2 according to the procedure described in the synthesis method of Compound No. A-1." When the compounds were synthesized according to the procedure described in the synthesis method of Compound No. A-6, TMAD or di-t-butyl azodicarboxylate (hereinafter abbreviated as "DBAB") was sometimes used instead of 40% DIAD.



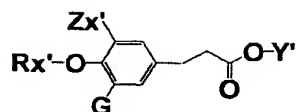


Table-A-1

Exp.	Rx'O	Y'	Zx'	G	Syn	LCMS		
						method	RTime	Mass
A-1	cPenMeO	Me	H	Br	A-1	C		341 (M <sup>+</sup> +1)
A-2	cHexMeO	Me	H	Br	A-2	C		354 (M <sup>+</sup> )
A-3	iBuO	Me	H	Br	A-2	A	5.34	N.D
A-4	2EtBuO	Me	H	Br	A-2			
A-5	cPenO	Me	H	Br	A-5	C		326 (M <sup>+</sup> )
A-6	cHexO	Me	H	Br	A-6	C		340 (M <sup>+</sup> )
A-7	cHepO	Me	H	Br	A-6			
A-8	BnO	Me	H	Br	A-2			
A-9	1PhEtO	Me	H	Br	A-2			
A-10	2FBnO	Me	H	Br	A-2			
A-11	4FBnO	Me	H	Br	A-2			
A-12	2ClBnO	Me	H	Br	A-2			
A-13	4ClBnO	Me	H	Br	A-2	A	4.85	N.D
A-14	4MeBnO	Me	H	Br	A-2			
A-15	4CF3BnO	Me	H	Br	A-2			
A-16	2(4DMAPh)EtO	Me	H	Br	A-6			
A-17	2(PhO)EtO	Me	H	Br	A-6	A	5.04	N.D
A-18	1(2FPh)EtO	Me	H	Br	A-6			
A-19	1(4ClPh)EtO	Me	H	Br	A-6	A	4.82	N.D
A-20	cPenMeO	Me	Cl	Br	A-20	C		375 (M <sup>+</sup> +1)
A-21	cPenMeO	Me	F	Br	A-21			
A-22	cPenO	Me	F	Br	A-21	C		345 (M <sup>+</sup> +1)
A-23	cHexO	Me	F	Br	A-21			
A-24	cPenO	Et	Me	Br	A-24	A	5.82	N.D
A-25	cPenO	Et	OMe	Br	A-25			
A-26	cPenMeO	Me	NO2	Br	A-26	C		340 (M <sup>+</sup> +1)
A-27	cHexMeO	Me	NO2	Br	A-26			
A-28	cPenO	Me	NO2	Br	A-26	C		372 (M <sup>+</sup> +1)
A-29	cHexO	Me	NO2	Br	A-26			
A-30	2-IndanO	Me	NO2	Br	A-26	A	5.03	N.D
A-31	PhO	Me	H	Br	A-31	A	5.15	N.D
A-32	4ClPhO	Me	H	Br	A-31	A	5.47	N.D
A-33	4MeOPhO	Me	H	Br	A-31	A	5.02	N.D

[Examples B-1 to B-119]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification including the examples described above are shown in Table B-1 to Table B-3.

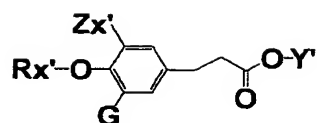


Table-B-1

Exp.	Rx'O	Y'	Zx'	G	Syn	LCMS		
						method	RTime	Mass
B-1	nPrO	Me	H	Br	A-2	C		279 (M <sup>+</sup> )
B-2	iPrO	Me	H	Br	A-2			
B-3	sBuO	Me	H	Br	A-6			
B-4	iPenO	Me	H	Br	A-6			
B-5	1,3DMeBuO	Me	H	Br	A-6			
B-6	2MeBuO	Me	H	Br	A-6			
B-7		Me	H	Br	A-6			
B-8		Me	H	Br	A-6			
B-9	2,3DMeBuO	Me	H	Br	A-6			
B-10	cPenO	Me	H	Cl	A-6	C		361 (M <sup>+</sup> +1)
B-11	trans2Me,cPenO	Me	H	Br	A-6			
B-12	3Me,cPenO	Me	H	Br	A-6			
B-13	trans2Me,cHexO	Me	H	Br	A-6			
B-14	cis2Me,cHexO	Me	H	Br	A-6			
B-15	3Me,cHexO	Me	H	Br	A-6	C		354 (M <sup>+</sup> +1)
B-16	4Me,cHexO	Me	H	Br	A-6			
B-17	2,3DMe,cHexO	Me	H	Br	A-6			
B-18	3,4DMe,cHexO	Me	H	Br	A-6	C		368 (M <sup>+</sup> +1)
B-19	3,5DMe,cHexO	Me	H	Br	A-6			
B-20		Me	H	Br	A-6			
B-21		Me	H	Br	A-6			
B-22		Me	H	Br	A-6			
B-23	1PhPrO	Me	H	Br	A-6			
B-24	(S)1PhPrO	Me	H	Br	A-6			
B-25	BenzhydrylO	Me	H	Br	A-6			
B-26		Me	H	Br	A-6	C		391 (M <sup>+</sup> +1)
B-27		Me	H	Br	A-6			
B-28	2Ph,1MeEtO	Me	H	Br	A-6			
B-29	2Ph,2MeEtO	Me	H	Br	A-6			
B-30	2(2FPh),1MeEtO	Me	H	Br	A-6			
B-31	2(3CF <sub>3</sub> Ph),1MeEtO	Me	H	Br	A-6			
B-32	3PhBuO	Me	H	Br	A-6			
B-33	5OMe-2-IndanO	Me	H	Br	A-6			
B-34	5,6D(OMe)-2-IndanO	Me	H	Br	A-6			
B-35	5F-2-IndaneO	Me	H	Br	A-6			
B-36	1-IndaneO	Me	H	Br	A-6			
B-37		Me	H	Br	A-6			

Table-B-2

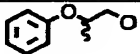
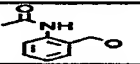








B-38		Me	H	Br	A-6			
B-39	3FBnO	Me	H	Br	A-6			
B-40	2MeBnO	Me	H	Br	A-6	C		363 (M <sup>+</sup> +1)
B-41	3MeBnO	Me	H	Br	A-6			
B-42	3,5DMeBnO	Me	H	Br	A-6			
B-43	4tBuBnO	Me	H	Br	A-6			
B-44	2CF <sub>3</sub> BnO	Me	H	Br	A-6			
B-45	4CF <sub>3</sub> BnO	Me	H	Br	A-6			
B-46	3(CF <sub>3</sub> O)BnO	Me	H	Br	A-6			
B-47	4(CF <sub>3</sub> O)BnO	Me	H	Br	A-6			
B-48	4(nBuO)BnO	Me	H	Br	A-6			
B-49		Me	H	Br	A-6	C		406 (M <sup>+</sup> +1)
B-50	3,4DFBnO	Me	H	Br	A-6			
B-51	2,4DFBnO	Me	H	Br	A-6			
B-52	4Br,2FBnO	Me	H	Br	A-6			
B-53	2,4DCIBnO	Me	H	Br	A-6			
B-54	3,4DCIBnO	Me	H	Br	A-6			
B-55	2,3DCIBnO	Me	H	Br	A-6			
B-56	2,6DCIBnO	Me	H	Br	A-6			
B-57	3,5DCIBnO	Me	H	Br	A-6			
B-58	2-NapMeO	Me	H	Br	A-6	C		399 (M <sup>+</sup> +1)
B-59	1-NapMeO	Me	H	Br	A-6			
B-60		Me	H	Br	A-6			
B-61		Me	H	Br	A-6			
B-62		Me	H	Br	A-6	C		339 (M <sup>+</sup> +1)
B-63	2PhBnO	Me	H	Br	A-6			
B-64	4PhBnO	Me	H	Br	A-6			
B-65	2PhEtO	Me	H	Br	A-6			
B-66	2(2MePh)EtO	Me	H	Br	A-6			
B-67	2(3MePh)EtO	Me	H	Br	A-6			
B-68	2(4MePh)EtO	Me	H	Br	A-6			
B-69	2(3FPh)EtO	Me	H	Br	A-6			
B-70	2(3ClPh)EtO	Me	H	Br	A-6			
B-71	2(2CF <sub>3</sub> Ph)EtO	Me	H	Br	A-6			
B-72	2(4CF <sub>3</sub> Ph)EtO	Me	H	Br	A-6			
B-73	2(2OMePh)EtO	Me	H	Br	A-6			
B-74	2(2-Nap)EtO	Me	H	Br	A-6	C		413 (M <sup>+</sup> +1)
B-75	2(3-Ind)EtO	Me	H	Br	A-6			
B-76		Me	H	Br	A-6			
B-77	2(PhO)EtO	Me	H	Br	A-6			
B-78	2(2ClPhO)EtO	Me	H	Br	A-6			
B-79	2(4ClPhO)EtO	Me	H	Br	A-6			

Table-B-3

B-80		Me	H	Br	A-6	C		407 (M <sup>+</sup> +1)
B-81		Me	H	Br	A-6			
B-82		Me	H	Br	A-6			
B-83		Me	H	Br	A-6			
B-84	2(PhS)EtO	Me	H	Br	A-6	C		379 (M <sup>+</sup> +1)
B-85	2-BztO	Me	H	Br	A-6			
B-86	(6OMe-2-Bzt)O	Me	H	Br	A-6			
B-87	cPenO	Me	Cl	Br	A-20			
B-88	1(4FPh)EtO	Me	Cl	Br	A-20			
B-89	1PhEtO	Me	F	Br	A-21			
B-90	1(4FPh)EtO	Me	F	Br	A-21			
B-91	1PhEtO	Et	Me	Br	A-24			
B-92	1(4FPh)EtO	Et	Me	Br	A-24			
B-93	1PhEtO	Me	OMe	Br	A-25			
B-94	1(4FPh)EtO	Me	OMe	Br	A-25			
B-95	BnO	Me	NO <sub>2</sub>	Br	A-26			
B-96	2-IndanO	Me	NO <sub>2</sub>	Br	A-26	A	4.44	N.D
B-97	5OMe-2-IndanO	Me	NO <sub>2</sub>	Br	A-26			
B-98	4CF <sub>3</sub> BnO	Me	NO <sub>2</sub>	Br	A-26			
B-99	cPenO	Me	NH <sub>2</sub>	Br	B-99	C		342 (M <sup>+</sup> +1)
B-100	BnO	Me	NH <sub>2</sub>	Br	B-99			
B-101	1PhEtO	Me	NH <sub>2</sub>	Br	B-99			
B-102	5OMe-2-IndanO	Me	NH <sub>2</sub>	Br	B-99			
B-103	BnO	Me	NHTFA	Br	B-103			
B-104	cPenO	Me	NHTFA	Br	B-103	C		438 (M <sup>+</sup> +1)
B-105	BnO	Me	NHMe	Br	B-105			
B-106	cPenO	Me	NHMe	Br	B-105	C		356 (M <sup>+</sup> +1)
B-107	1PhEtO	Me	NHMe	Br	B-105			
B-108	1(4FPh)EtO	Me	NHMe	Br	B-105			
B-109	BnO	Me	NMe <sub>2</sub>	Br	B-109			
B-110	cPenO	Me	NMe <sub>2</sub>	Br	B-109	C		370 (M <sup>+</sup> +1)
B-111	1PhEtO	Me	NMe <sub>2</sub>	Br	B-109			
B-112	1(4FPh)EtO	Me	NMe <sub>2</sub>	Br	B-109			
B-113	cPenO	Me	OH	Br	B-113	C		343 (M <sup>+</sup> +1)
B-114	cPenO	Me	OCOMe	Br	B-114			
B-115	1(4FPh)EtO	Me	OH	Br	B-113			
B-116	1(4FPh)EtO	Me	OCOMe	Br	B-114			
B-117	cPenMeO	Me	Br	Br	B-117			
B-118	cPenO	Me	Br	Br	B-117	A	5.98	N.D
B-119	1(4FPh)EtO	Me	Br	Br	B-117			

[Example C-1]

Synthesis of 3-bromo-4-cyclohexylmethyloxybenzaldehyde (Intermediate 25)

According to the procedure described in the synthesis method of Compound

No. A-2 provided that the purification was performed by flash column chromatography (hexane: isopropyl ether = 5:1), 3-bromo-4-hydroxybenzaldehyde (17.4 g), potassium carbonate (23.9 g) and bromomethylcyclohexane (36.2 ml) were reacted and treated to obtain the title compound (Intermediate 25, 18.7 g).

Synthesis of 4-cyclohexylmethoxy-3-(naphthalen-2-yl)benzaldehyde (Compound No. C-1)

A solution of 2-naphthaleneboronic acid (535 mg) in methanol (5.0 ml), Intermediate 25 (1.16 g), and 2 M aqueous sodium carbonate (0.9 ml) were added with toluene (10.0 ml) and tetrakis(triphenylphosphine)palladium(0) [hereinafter abbreviated as "(Ph<sub>3</sub>P)<sub>4</sub>Pd"] (116 mg, Nakarai Tecs), and stirred at 80°C for 17 hours. The reaction mixture was added with ethyl acetate (100 ml), and washed successively with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride and saturated brine. The organic layer was dried, and then the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography (hexane:ethyl acetate = 10:1) to obtain the title compound (Compound No. C-1, 345 mg).

[Example D-10]

Synthesis of 3-bromo-4-hydroxy-5-nitrobenzaldehyde (Intermediate 26)

A solution of 3-bromo-4-hydroxybenzaldehyde (6.30 g) in acetic acid (45 ml) was added dropwise with 70% nitric acid (5.85 ml) on a water bath, then added with sodium nitrite (62 mg), and further stirred for 2 hours. The reaction mixture was poured into ice water (300 ml), and precipitates were taken by filtration, and washed with water (50ml x 3). The precipitates were dried under reduced pressure for 24 hours to obtain the title compound (Intermediate 26, 5.88 g).

Synthesis of 3-bromo-4-cyclohexylmethoxy-5-nitrobenzaldehyde (Intermediate 27)

According to the procedure described in the synthesis method of Compound No. A-2 provided that the purification was performed by flash column

chromatography (hexane:ethyl acetate = 7:1), Intermediate 26 (5.5 g), potassium carbonate (3.94 g) and bromomethylcyclohexane (3.94 ml) were reacted and treated to obtain the title compound (Intermediate 27, 5.2 g).

Synthesis of 4-cyclohexylmethoxy-3-(naphthalen-2-yl)-5-nitrobenzaldehyde  
(Compound No. D-10)

According to the procedure described in the synthesis method of Compound No. C-1 with the modifications that the reaction was carried out for 15 hours at 80°C, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 7:1), Intermediate 27 (2.65 g), 2-naphthaleneboronic acid (3.01 g), 2 M aqueous sodium carbonate (7.5 ml) and (Ph<sub>3</sub>P)<sub>4</sub>Pd (960 mg) were reacted and treated to obtain the title compound (Compound No. D-10, 2.96 g).

[Examples C-1 to C-8]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification including the examples described above are shown in Table C-1.

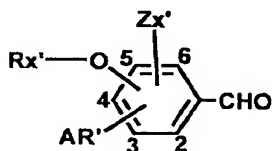


Table-C-1

Exp.	Rx'O	Rx'O Position	Zx'	Zx' Position	AR'	AR' Position	Syn	LCMS		
								method	RTtime	Mass
C-1	cHexMeO	4	H	-	2-Nap	5	C-1			
C-2	cHexMeO	4	H	-	1-Nap	5	C-1			
C-3	cHexMeO	4	H	-	2OMe-6-Nap	5	C-1	C		374(M <sup>+</sup> )
C-4	cHexMeO	4	H	-	5-Ind	5	C-1			
C-5	cPenMeO	4	H	-	2-Nap	5	C-1			
C-6	cPenMeO	4	H	-	5-Ind	5	C-1			
C-7	cPenO	4	H	-	2-Nap	5	C-1	C		316(M <sup>+</sup> )
C-8	cPenO	4	H	-	5-Ind	5	C-1	C		305(M <sup>+</sup> )

[Examples D-1 to D-29]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification including the examples described above are shown in Table-D-1.

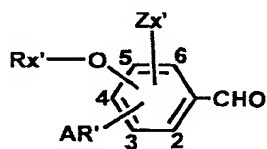


Table-D-1

Exp.	Rx'O	Rx'O Position	Zx'	Zx' Position	AR'	AR' Position	Syn	LCMS		
								method	RTTime	Mass
D-1	cHexMeO	4	H	—	2-BT	3	C-1	C		350 (M <sup>+</sup> )
D-2	cHexMeO	4	H	—	2-BF	3	C-1			
D-3	cHexMeO	4	H	—	1Me-5-Ind	3	C-1	C		316(M <sup>+</sup> )
D-4	cHexMeO	4	H	—	5-1Hldz	3	C-1			
D-5	cHexMeO	4	H	—	1Me-5-1Hldz	3	C-1			
D-6	2(2FPh)EtO	4	H	—	2-Nap	3	C-1			
D-7	2(2FPh)EtO	4	H	—	5-Ind	3	C-1			
D-8	2-IndanO	4	H	—	5-Ind	3	C-1			
D-9	2-IndanO	4	H	—	5-1Hldz	3	C-1			
D-10	cPenMeO	4	NO <sub>2</sub>	5	2-Nap	3	D-10	C		330 (M <sup>+</sup> +1)
D-11	cPenMeO	4	NO <sub>2</sub>	5	5-Ind	3	D-10			
D-12	cHexMeO	4	NO <sub>2</sub>	5	2-Nap	3	D-10			
D-13	cHexMeO	4	NO <sub>2</sub>	5	2-BF	3	D-10			
D-14	cPenO	4	NO <sub>2</sub>	5	2-Nap	3	D-10			
D-15	cPenO	4	NO <sub>2</sub>	5	5-Ind	3	D-10	C		350(M <sup>+</sup> )
D-16	2(2FPh)EtO	4	NO <sub>2</sub>	5	2-Nap	3	D-10			
D-17	2(2FPh)EtO	4	NO <sub>2</sub>	5	5-Ind	3	D-10			
D-18	2-IndanO	4	NO <sub>2</sub>	5	5-Ind	3	D-10			
D-19	2-IndanO	4	NO <sub>2</sub>	5	1Me-5-1Hldz	3	D-10	A	3.85	414 (M <sup>+</sup> +1)
D-20	cPenO	2	H	—	2-Nap	5	C-1	C		316(M <sup>+</sup> )
D-21	cPenO	2	H	—	5-Ind	5	C-1	C		305(M <sup>+</sup> )
D-22	cPenO	3	H	—	2-Nap	5	C-1			
D-23	cPenO	3	H	—	5-Ind	5	C-1			
D-24	cPenO	5	H	—	2-Nap	2	C-1			
D-25	cPenO	5	H	—	5-Ind	2	C-1			
D-26	cPenO	4	H	—	2-Nap	2	C-1			
D-27	cPenO	4	H	—	5-Ind	2	C-1			
D-28	cPenO	3	H	—	2-Nap	2	C-1			
D-29	cPenO	3	H	—	5-Ind	2	C-1			

#### [Example E-1]

#### Synthesis of 5-bromo-2-cyclopentylmethoxypyridine (Intermediate 28)

A solution of potassium t-butoxide (550.6 mg, WAKO) in dehydrated THF (10 ml) was added with cyclopentane methanol (450  $\mu$ l), and then added with a solution of 2,5-dibromopyridine (982.8 mg, TCI) in dehydrated THF (15 ml) under

ice cooling. The reaction mixture was stirred for 30 minutes, then warmed to room temperature, and stirred for 11 hours. The reaction mixture was added with water (100 ml) and ethyl acetate (60 ml) for extraction. The organic layer was washed successively with saturated aqueous sodium hydrogencarbonate and saturated brine sequentially, and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 15:1) to obtain the title compound (Intermediate 28, 896 mg).  
Synthesis of 2-cyclopentylmethyloxypyridine-5-carbaldehyde (Intermediate 29)

A solution of Intermediate 28 (895 mg) in anhydrous THF (10 ml) was added dropwise with a 1.6 M solution of n-butyllithium in hexane (2.70 ml, Ald) over 5 minutes with cooling at -78°C under argon gas atmosphere, and stirred for 20 minutes. This reaction mixture was added with dehydrated DMF (330  $\mu$ l, WAKO) over 3 minutes, stirred for 30 minutes, then warmed to room temperature, and further stirred for 1 hour. The reaction mixture was added with water (10 ml), and extracted with ethyl acetate (30ml x 3). The organic layer was washed with saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 10:1) to obtain the title compound (Intermediate 29, 1.04 g).  
Synthesis of ethyl 3-(2-cyclopentylmethyloxypyridin-5-yl)acrylate (Intermediate 30)

According to the procedure described in the synthesis method of Intermediate 7 with the modification that the reaction was carried out for 1 hour, Intermediate 29 (450 mg), ethyl diethylphosphonoacetate (530  $\mu$ l) and 60% sodium hydride (120 mg) were reacted and treated to obtain the title compound (Intermediate 30, 394 mg).

Synthesis of ethyl 3-(2-cyclopentylmethyloxypyridine-5-yl)propionate (Intermediate 31)

According to the procedure described in the synthesis method of



Intermediate 8 with the modifications that the reaction was carried out for 1 hour, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 15:1), Intermediate 30 (392 mg) and 10% palladium/carbon (30 mg) were reacted and treated to obtain the title compound (Intermediate 31, 246 mg).

Synthesis of ethyl 3-(3-bromo-2-cyclopentylmethoxy)pyridin-5-yl)propionate (Compound No. E-1)

A solution of Intermediate 31 (5.20 g) in acetonitrile (50 ml) was warmed to 35°C, added dropwise with bromine (1.1 ml, WAKO), then added with NBS (3.72 g), and stirred at room temperature for 2 hours. The reaction mixture was concentrated under reduced pressure, then added with ethyl acetate (200 ml), and washed successively with saturated aqueous ammonium chloride, 5% aqueous sodium sulfite, saturated aqueous sodium hydrogencarbonate and saturated brine. The organic layer was dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 10:1) to obtain the title compound (Compound No. E-1, 6.51 g).

[Example E-7]

Synthesis of 2-benzyloxy-5-bromopyridine (Intermediate 32)

According to the procedure described in the synthesis method of Intermediate 28 provided that the reaction was carried out for 1 hour, potassium *t*-butoxide (3.13 g), benzyl alcohol (3.10 ml) and 2,5-dibromopyridine (4.79 g) were reacted and treated to obtain the title compound (Intermediate 32, 5.36 g).

Synthesis of 2-benzyloxypyridine-5-carbaldehyde (Intermediate 33)

According to the procedure described in the synthesis method of Intermediate 29, Intermediate 32 (5.10 g), a 1.6M solution of *n*-butyllithium in hexane (15.5 ml) and dehydrated DMF (1.9 ml) were reacted and treated to obtain the title compound (Intermediate 33, 2.75 g).

#### Synthesis of ethyl 3-(2-benzyloxypyridin-5-yl)acrylate (Intermediate 34)

According to the procedure described in the synthesis method of Intermediate 7, Intermediate 33 (2.74 g), ethyl diethylphosphonoacetate (3.12 ml) and 60% sodium hydride (635 mg) were reacted and treated to obtain the title compound (Intermediate 34, 2.12 g).

#### Synthesis of ethyl 3-(2-hydroxypyridin-5-yl)propionate (Intermediate 35)

According to the procedure described in the synthesis method of Intermediate 8 provided that the reaction was carried out for 2.5 hours, Intermediate 54 (2.12 g) and 10% palladium/carbon (120 mg) were reacted and treated to obtain the title compound (Intermediate 35, 1.26 g).

#### Synthesis of ethyl 3-(3-bromo-2-hydroxypyridin-5-yl)propionate (Intermediate 36)

According to the procedure described in the synthesis method of Compound No. E-1 with the modifications that the reaction was carried out for 2.5 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 1:2), Intermediate 35 (1.23 g), bromine (340  $\mu$ l) and NBS (1.19 g) were reacted and treated to obtain the title compound (Compound No. 36, 1.42 g).

#### Synthesis of ethyl 3-[5-bromo-6-[(S)-1-phenylethyloxy]pyridin-3-yl]propionate (Compound No. E-7)

According to the procedure described in the synthesis method of Compound No. A-6 with the modifications that the reaction was carried out for 11 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 4:1), Intermediate 36 (137 mg),  $\text{Ph}_3\text{P}$  (273 mg), (R)-1-phenylethanol (150  $\mu$ l, TCI) and 40% DIAD (400  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. E-7, 167 mg).

#### [Example E-13]

#### Synthesis of ethyl 3-(5-bromo-6-(4-trifluoromethylbenzyloxy)pyridin-3-yl)propionate (Compound No. E-13)

A solution of Intermediate 36 (71.5 mg) in chloroform (7 ml) was added with 4-trifluoromethylbenzyl bromide (109.2 mg, TCI) and silver carbonate (120 mg, WAKO), and stirred at room temperature for 11 hours under light shielding. The reaction mixture was filtered, and the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 6:1) to obtain the title compound (Compound No. E-13, 114 mg).

[Example E-1 to 16]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification including the examples described above are shown in Table E-1.

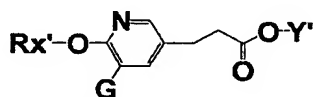


Table-E-1

Exp.	Rx'O	Y'	G	Syn	LCMS		
					method	RTime	Mass
E-1	cPenMeO	Et	Br	E-1	A	5.98	356(M <sup>+</sup> )
E-2	cHexMeO	Et	Br	E-1			
E-3	iBuO	Et	Br	E-1	A	5.57	N.D
E-4	2EtBuO	Et	Br	E-1			
E-5	cPenO	Et	Br	E-1	A	5.62	342 (M <sup>+</sup> )
E-6	cHexO	Et	Br	E-1			
E-7	(R)1PhEtO	Et	Br	E-7	A	5.60	N.D
E-8	2(4DMAPh)EtO	Et	Br	E-7			
E-9	2(2FPh)EtO	Et	Br	E-7			
E-10	2(3FPh)EtO	Et	Br	E-7			
E-11	2(4ClPh)EtO	Et	Br	E-7			
E-12	2(PhO)EtO	Et	Br	E-7			
E-13	4CF <sub>3</sub> BnO	Et	Br	E-13	A	5.78	432 (M <sup>+</sup> )
E-14	2MeBnO	Et	Br	E-13			
E-15	2ClBnO	Et	Br	E-13			
E-16	1(4FPh)EtO	Et	Br	E-7			

[Example F-1]

Synthesis of 4-(3-bromo-4-methoxyphenyl)butyric acid (Intermediate 37)

According to the procedure described in the synthesis method of Compound No. A-1 provided that the reaction was carried out under ice cooling for 30 minutes and for 20 hours at room temperature, 4-(4-methoxyphenyl)butyric acid (11.64 g, Ald) and NBS (11.21 g) were reacted and treated to obtain the title compound (Intermediate 37, 16.30 g).

#### Synthesis of methyl 4-(3-bromo-4-hydroxyphenyl)butyrate (Intermediate 38)

According to the procedure described in the synthesis method of Intermediate 4, Intermediate 37 (12.51 g) and a 1 M solution of boron tribromide in methylene chloride (100 ml) were reacted and treated, and the obtained residue was reacted with thionyl chloride (8.4 ml) in methanol and treated according to the procedure described in the synthesis method of Intermediate 1 to obtain the title compound (Intermediate 38, 10.48 g).

#### Synthesis of methyl 4-(3-bromo-4-cyclopentylmethoxyphenyl)butyrate (Compound No. F-1)

According to the procedure described in the synthesis method of Compound No. A-6 provided that the purification was performed by column chromatography (Quad, hexane:isopropyl alcohol = 10:1), Intermediate 38 (2.72 g),  $\text{Ph}_3\text{P}$  (7.86 g), cyclopentane methanol (3.24 ml) and 40% DIAD (14.2 ml) were reacted and treated to obtain the title compound (Compound No. F-1, 3.33 g).

#### [Examples F-1 to F-4]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification including the examples described above are shown in Table-F-1.

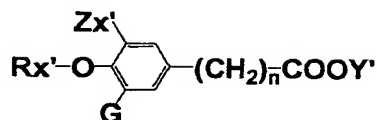


Table-F-1

Exp.	Rx'O	Y'	Zx'	G	n	Syn	LCMS		
							method	RTime	Mass
F-1	cPenMeO	Me	H	Br	3	F-1	C		354(M <sup>+</sup> )
F-2	cPenO	Me	H	Br	3	F-1			
F-3	cHexO	Me	H	Br	3	F-1	C		354(M <sup>+</sup> )
F-4	1(4FPh)EtO	Me	H	Br	3	F-1			

[Example G-1]

Synthesis of methyl 3-[4-methoxy-3-(naphthalen-2-yl)phenyl]propionate

(Intermediate 39)

According to the procedure described in the synthesis method of Compound No. C-1 with the modifications that the reaction was carried out for 2 hours, and the purification was performed by flash column chromatography (hexane:isopropyl ether = 8:1), Intermediate 20 (460 mg), 2-naphthaleneboronic acid (886 mg), 2 M aqueous sodium carbonate (1.6 ml) and (Ph<sub>3</sub>P)<sub>4</sub>Pd (298 mg) were reacted and treated to obtain the title compound (Intermediate 39, 580 mg).

Synthesis of 3-[4-methoxy-3-(naphthalen-2-yl)phenyl]propionic acid (Intermediate 40)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 2 hours, Intermediate 39 (773 mg) and 2 N aqueous sodium hydroxide (2.3 ml) were reacted and treated to obtain the title compound (Intermediate 40, 674 mg).

Synthesis of methyl 3-[4-hydroxy-3-(naphthalen-2-yl)phenyl]propionate

(Intermediate 41)

According to the procedure described in the synthesis method of Intermediate 10, pyridine (5 ml), concentrated hydrochloric acid (5 ml), and Intermediate 40 (551 mg) were reacted and treated to obtain crude powder

substance. This substance was reacted with thionyl chloride (282  $\mu$ l) in methanol and treated according to the procedure described in the synthesis method of Intermediate 1 to obtain the title compound (Intermediate 41, 531 mg).

Synthesis of methyl 3-[4-cyclopentyloxy-3-(naphthalen-2-yl)phenyl]propionate  
(Compound No. G-1)

According to the procedure described in the synthesis method of Compound No. A-6 with the modifications that the reaction was carried out for 15 hours, and the purification was performed by flash column chromatography (hexane:isopropyl ether = 6:1), Intermediate 41 (100 mg),  $\text{Ph}_3\text{P}$  (262 mg), cyclopentanol (91  $\mu$ l, TCI) and 40% DIAD (473  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. G-1, 120 mg).

[Example G-2]

Synthesis of 3-[4-cyclopentyloxy-3-(naphthalen-2-yl)phenyl]propionic acid  
(Compound No. G-2)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 4 hours, Compound No. G-1 (115 mg), and 2 N aqueous sodium hydroxide (0.75 ml) were reacted and treated to obtain the title compound (Compound No. G-2, 108 mg).

[Example G-3]

Synthesis of methyl 3-[4-cyclopentyloxy-3-(1H-indol-5-yl)phenyl]propionate  
(Compound No. G-3)

According to the procedure described in the synthesis method of Compound No. C-1 with the modifications that the reaction was carried out for 3 hours, and the purification was performed by flash column chromatography (hexane:ethyl acetate = 4:1), Compound No. A-5 (833 mg), 5-indoleboronic acid (657 mg), 2 M aqueous sodium carbonate (2.4 ml) and  $(\text{Ph}_3\text{P})_4\text{Pd}$  (233 mg) were reacted and treated to obtain the title compound (Compound No. G-3, 900 mg).

[Example G-4]

Synthesis of 3-[4-cyclopentyloxy-3-(1H-indole-5-yl)phenyl]propionic acid (Compound No. G-4)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 2 hours, Compound No. G-3 (144 mg) and 2 N aqueous sodium hydroxide (420  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. G-4, 127 mg).

[Example G-9]

Synthesis of methyl 3-[4-benzyloxy-5-(1-methyl-1H-indazol-5-yl)phenyl]propionate (Compound No. G-9)

According to the procedure described in the synthesis method of Compound No. C-1 with the modifications that the reaction was carried out at 80°C for 6 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 4:1), Compound No. A-8 (349 mg), 1-methyl-1H-indazole-5-boronic acid (283 mg), 2 M aqueous sodium carbonate (0.9 ml) and (Ph<sub>3</sub>P)<sub>4</sub>Pd (94.3 mg) were reacted and treated to obtain the title compound (Compound No. G-9, 370 mg).

[Example G-10]

Synthesis of 3-[4-benzyloxy-5-(1-methyl-1H-indazol-5-yl)phenyl]propionic acid (Compound No. G-10)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 4 hours, Compound No. G-9 (80 mg) and 2 N aqueous sodium hydroxide (0.20 ml) were reacted and treated to obtain the title compound (Compound No. G-10, 71 mg).

Synthesis of methyl 3-[4-hydroxy-5-(1-methyl-1H-indazol-5-yl)phenyl]propionate (Intermediate 42)

A solution of Compound No. G-9 (314 mg) in a mixture of ethyl acetate (3 ml) and methanol (3 ml) was added with 10% palladium/carbon (12 mg), and stirred

at room temperature for 16 hours under hydrogen atmosphere. The reaction mixture was filtered, and the solvent of the filtrate was evaporated under reduced pressure to obtain the title compound (Intermediate 48, 288 mg).

[Example G-23]

Synthesis of methyl 3-(3-bromo-4-t-butyldimethylsilyloxyphenyl)propionate (Intermediate 43)

According to the procedure described in the synthesis method of Intermediate 16 provided that the reaction was carried out for 16 hours, Intermediate 5 (5.18 g), imidazole (2.04 g) and t-butyldimethylsilyl chloride (4.52 g) were reacted and treated to obtain the title compound (Intermediate 43, 8.42 g).

Synthesis of methyl 3-[4-(t-butyldimethylsilyloxy-3-(1H-indol-5-yl)phenyl)propionate (Intermediate 44)

According to the procedure described in the synthesis method of Compound No. C-1 with the modifications that reaction was performed for 12.5 hours, and the purification was performed by flash column chromatography (hexane:ethyl acetate = 9:1), 5-indoleboronic acid (4.83 g), Intermediate 34 (7.46 g), 2 M aqueous sodium carbonate (18 ml) and (Ph<sub>3</sub>P)<sub>4</sub>Pd (1.62 g) were reacted and treated to obtain the title compound (Intermediate 44, 5.04 g).

Synthesis of methyl 3-[4-hydroxy-3-(1H-indol-5-yl)phenyl]propionate (Intermediate 45)

According to the procedure described in the synthesis method of Intermediate 19 with the modifications that the reaction was carried out for 2 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 3:1), Intermediate 35 (5.04 g), acetic acid (2.8 ml) and a 1 M solution of tetrabutylammonium fluoride in THF (49 ml, TCI) were reacted and treated to obtain the title compound (Intermediate 45, 3.13 g).

Synthesis of methyl 3-[3-(1H-indol-5-yl)-4-(4-methylphenylmethoxy)phenyl]-



propionate (Compound No. G-23)

According to the procedure described in the synthesis method of Compound No. A-2 with the modifications that the reaction was carried out for 15 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 5:1), Intermediate 45 (80 mg), potassium carbonate (114 mg) and 4-methylbenzyl bromide (54  $\mu$ l, TCI) were reacted and treated to obtain the title compound (Compound No. G-23, 104 mg).

[Example G-24]

Synthesis of 3-[3-(1H-indol-5-yl)-4-(4-methylphenylmethoxy)phenyl]propionic acid (Compound No. G-24)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 3 hours, Compound No. G-23 (99 mg) and 2 N aqueous sodium hydroxide (500  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. G-24, 84 mg).

[Example G-106]

Synthesis of N-[2-(t-butyldiphenylsilyloxy)ethyl]aniline (Intermediate 46)

A solution of 2-anilinoethanol (5.82 g, TCI) in anhydrous DMF (50 ml) was added with imidazole (3.23 g, TCI), added dropwise with a solution of t-butyldiphenylsilyl chloride (12.48 g, TCI) in DMF (50 ml) under ice cooling, stirred for 30 minutes, then warmed to room temperature, and further stirred for 3.5 hours. The reaction mixture was added with water (100 ml), and extracted with ethyl acetate (100 ml). The organic layer was washed successively with water and saturated brine, and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography (hexane:ethyl acetate = 9:1) to obtain the title compound (Intermediate 46, 15.61 g).

Synthesis of N-benzyl-N-[2-(t-butyldiphenylsilyloxy)ethyl]aniline (Intermediate 47)

According to the procedure described in the synthesis method of Compound

No. A-2 with the modifications that the reaction was carried out for 15 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 5:1), Intermediate 46 (15.60 g), potassium carbonate (8.91 g) and benzyl bromide (6.05 ml, TCI) were reacted and treated to obtain the title compound (Intermediate 47, 19.23 g).

Synthesis of 2-(N-benzyl-N-phenylamino)ethanol (Intermediate 48)

According to the procedure described in the synthesis method of Intermediate 9 with the modifications that the reaction was carried out for 1 hour, and the purification was performed by flash column chromatography (hexane:ethyl acetate = 5:1), Intermediate 47 (19.22 g) and a 1 M solution of tetrabutylammonium fluoride in THF (86 ml) were reacted and treated to obtain the title compound (Intermediate 48, 9.06 g).

Synthesis of methyl 3-{4-[2-(N-benzyl-N-phenylamino)ethoxy]-3-(naphthalen-2-yl)phenyl}propionate (Compound No. G-106)

According to the procedure described in the synthesis method of Compound No. A-6 with the modifications that the reaction was carried out for 15 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 7:1), Intermediate 41 (1.26 g), Ph<sub>3</sub>P (1.34 g), Intermediate 48 (1.01 g) and DBAB (1.18 g) instead of 40% DIAD were reacted and treated to obtain the title compound (Compound No. G-106, 1.39 g).

[Example G-107]

Synthesis of methyl 3-{3-(naphthalen-2-yl)-4-[2-(N-phenylamino)ethoxy]phenyl}propionate (Compound No. G-107)

A solution of Compound No. G-106 (1.39 g) in a mixture of THF (10 ml) and methanol (20 ml) was added with concentrated hydrochloric acid (75  $\mu$ l, WAKO) and 10% palladium/carbon (142 mg), and stirred at room temperature for 3 hours under hydrogen gas atmosphere. The reaction mixture was filtered, and the

solvent of the filtrate was evaporated under reduced pressure to obtain the title compound (Compound No. G-107, 842 mg).

[Example G-108]

Synthesis of 3-{3-(naphthalen-2-yl)-4-[2-(phenylamino)ethyloxy]phenyl}propionic acid (Compound No. G-108)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 2 hours, Compound No. G-107 (46 mg) and 2 N aqueous sodium hydroxide (0.25 ml) were reacted and treated to obtain the title compound (Compound No. G-108, 41 mg).

[Examples G-1 to G-121]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification including the examples described above are shown in Table-G-1 to Table-G-4.

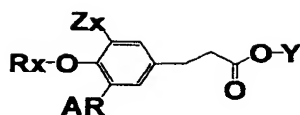


Table-G-1




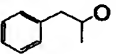
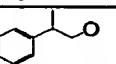
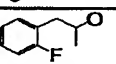
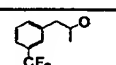
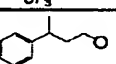
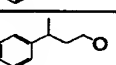
Exp.	RxO	Y	Zx	AR	Syn	LCMS		
						method	RTime	Mass
G-1	cPenMeO	Me	H	2-Nap	G-1	C		388(M <sup>+</sup> )
G-2	cPenMeO	H	H	2-Nap	G-2	C		375 (M <sup>+</sup> +1)
G-3	cPenMeO	Me	H	5-Ind	G-3			
G-4	cPenMeO	H	H	5-Ind	G-4	C		363 (M <sup>+</sup> )
G-5	cPenMeO	Me	H	1Me-5-Ind	G-3			
G-6	cPenMeO	H	H	1Me-5-Ind	G-4	A		391 (M <sup>+</sup> +1)
G-7	cPenMeO	Me	H	5-1HIdz	G-3			
G-8	cPenMeO	H	H	5-1HIdz	G-4			
G-9	BnO	Me	H	1Me-5-1HIdz	G-9			
G-10	BnO	H	H	1Me-5-1HIdz	G-10			
G-11	cPenMeO	Me	H	1Me-5-1HIdz	G-3			
G-12	cPenMeO	H	H	1Me-5-1HIdz	G-4			
G-13	2EtBuO	H	H	2-Nap	G-1,G-2	A		377 (M <sup>+</sup> +1)
G-14	2EtBuO	H	H	5-Ind	G-3,G-4			
G-15	4Me,cHexO	H	H	2-Nap	G-1,G-2			
G-16	4Me,cHexO	H	H	5-Ind	G-3,G-4	D	5.46	378 (M <sup>+</sup> +1)
G-17		H	H	2-Nap	G-1,G-2			
G-18		H	H	5-Ind	G-3,G-4			
G-19	cHepO	H	H	5-Ind	G-3,G-4			
G-20	3PhPrO	H	H	2-Nap	G-1,G-2			
G-21	4PhBuO	H	H	5-Ind	G-3,G-4			
G-22		H	H	2-Nap	G-1,G-2	D	5.40	414 (M <sup>+</sup> +1)
G-23	4MeBnO	Me	H	5-Ind	G-23			
G-24	4MeBnO	H	H	5-Ind	G-24			
G-25	2(4MePh)EtO	H	H	2-Nap	G-1,G-2			
G-26	2(4MePh)EtO	H	H	5-Ind	G-1,G-2			
G-27	4ClBnO	H	H	2-Nap	G-23,G-24			
G-28	4CF <sub>3</sub> BnO	H	H	5-Ind	G-23,G-24			
G-29	3F,4(OMe)BnO	H	H	2-Nap	G-1,G-2			
G-30	3F,4(OMe)BnO	H	H	5-Ind	G-1,G-2			
G-31		H	H	2-Nap	G-1,G-2			
G-32		H	H	5-Ind	G-1,G-2			
G-33		H	H	2-Nap	G-1,G-2			
G-34		H	H	5-Ind	G-1,G-2			
G-35		H	H	2-Nap	G-1,G-2			
G-36		H	H	5-Ind	G-1,G-2			

Table-G-2

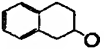
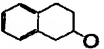
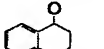
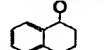
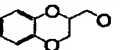
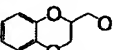
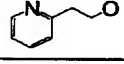
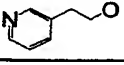
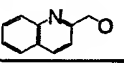
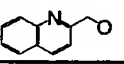
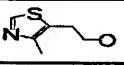
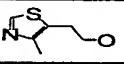
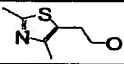
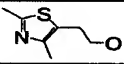
G-37	1IndanO	H	H	5-Ind	G-1,G-2	D	5.19	398 (M <sup>+</sup> +1)
G-38	2IndanO	H	H	2-Nap	G-1,G-2			
G-39	2IndanO	H	H	5-Ind	G-1,G-2			
G-40	5OMe-2-IndanO	H	H	2-Nap	G-1,G-2	C		439(M <sup>+</sup> +1)
G-41	5,6D(OMe)-2-IndanO	H	H	5-Ind	G-1,G-2	C		458(M <sup>+</sup> +1)
G-42	5F-2-IndanO	H	H	2-Nap	G-1,G-2			
G-43	5F-2-IndanO	H	H	5-Ind	G-1,G-2	C		416(M <sup>+</sup> +1)
G-44		H	H	2-Nap	G-1,G-2			
G-45		H	H	5-Ind	G-1,G-2	A	5.46	412 (M <sup>+</sup> +1)
G-46		H	H	2-Nap	G-1,G-2			
G-47		H	H	5-1HInd	G-1,G-2			
G-48	2(2MePh)EtO	H	H	2-Nap	G-1,G-2			
G-49	2(2MePh)EtO	H	H	5-Ind	G-1,G-2			
G-50	2(3FPh)EtO	H	H	2-Nap	G-1,G-2			
G-51	2(2ClPh)EtO	H	H	2-Nap	G-1,G-2			
G-52	2(3ClPh)EtO	H	H	5-Ind	G-1,G-2			
G-53	2(2CF <sub>3</sub> Ph)EtO	H	H	5-Ind	G-1,G-2			
G-54	4(CF <sub>3</sub> Ph)EtO	H	H	2-Nap	G-1,G-2			
G-55	2(2OMePh)EtO	H	H	2-Nap	G-1,G-2	C		427 (M <sup>+</sup> +1)
G-56	2(4OMePh)EtO	H	H	5-Ind	G-1,G-2			
G-57	2(1-NapEt)O	H	H	2-Nap	G-1,G-2			
G-58	2(2-Nap)EtO	H	H	2-Nap	G-1,G-2			
G-59	2(2-Nap)EtO	H	H	5-Ind	G-1,G-2	C		435 (M <sup>+</sup> )
G-60	2(4ClPh)EtO	H	H	2-Nap	G-1,G-2			
G-61		H	H	5-Ind	G-1,G-2	D	5.11	430 (M <sup>+</sup> +1)
G-62		H	H	1Me-5-1HIdz	G-1,G-2			
G-63	2(PhS)EtO	H	H	2-Nap	G-1,G-2	A		402 (M <sup>+</sup> +1)
G-64	2(PhS)EtO	H	H	5-Ind	G-1,G-2			
G-65	3PhPrO	H	H	5-Ind	G-1,G-2			
G-66	2ClBnO	H	H	2-Nap	G-1,G-2			
G-67	2BrBnO	H	H	5-Ind	G-1,G-2	C		450 (M <sup>+</sup> )
G-68	3,5DMeBnO	H	H	5-Ind	G-1,G-2			
G-69	4tBuBnO	H	H	2-Nap	G-1,G-2			
G-70	2CF <sub>3</sub> BnO	H	H	2-Nap	G-1,G-2			
G-71	4CF <sub>3</sub> BnO	H	H	5-Ind	G-1,G-2			
G-72	4nBuBnO	H	H	5-Ind	G-1,G-2			
G-73	3,5DClBnO	H	H	2-Nap	G-1,G-2			
G-74	2,3DClBnO	H	H	5-Ind	G-1,G-2			
G-75	2PhBnO	H	H	2-Nap	G-1,G-2			
G-76	4PhBnO	H	H	5-Ind	G-1,G-2	A		448 (M <sup>+</sup> +1)

Table-G-3

G-77		H	H	2-Nap	G-1,G-2			
G-78		H	H	5-Ind	G-1,G-2			
G-79		H	H	2-Nap	G-1,G-2			
G-80		H	H	5-Ind	G-1,G-2			
G-81		H	H	2-Nap	G-1,G-2	C		386 (M <sup>+</sup> +1)
G-82		H	H	5-Ind	G-1,G-2			
G-83		H	H	2-Nap	G-1,G-2			
G-84		H	H	5-Ind	G-1,G-2			
G-85		H	H	2-Nap	G-1,G-2			
G-86		H	H	5-Ind	G-1,G-2			
G-87		H	H	2-Nap	G-1,G-2			
G-88		H	H	5-Ind	G-1,G-2			
G-89		H	H	2-Nap	G-1,G-2			
G-90		H	H	5-Ind	G-1,G-2			
G-91		H	H	2-Nap	G-1,G-2			
G-92		H	H	5-Ind	G-1,G-2			
G-93		H	H	2-Nap	G-1,G-2			
G-94		H	H	2-Nap	G-1,G-2	C		384 (M <sup>+</sup> +1)
G-95		H	H	5-Ind	G-1,G-2			
G-96		H	H	2-Nap	G-1,G-2			
G-97		H	H	5-Ind	G-1,G-2			

Table-G-4

G-98		H	H	2-Nap	G-1,G-2			
G-99		H	H	5-Ind	G-1,G-2			
G-100		H	H	2-Nap	G-1,G-2			
G-101		H	H	5-Ind	G-1,G-2	C		423 (M <sup>+</sup> +1)
G-102		H	H	2-Nap	G-1,G-2			
G-103		H	H	5-Ind	G-1,G-2			
G-104		H	H	2-Nap	G-1,G-2			
G-105		H	H	5-Ind	G-1,G-2			
G-106	2(Ph,BnN)EtO	Me	H	2-Nap	G-106			
G-107	2(PhNH)EtO	Me	H	2-Nap	G-107			
G-108	2(PhNH)EtO	H	H	2-Nap	G-108	C		412(M <sup>+</sup> +1)
G-109	2(PhNH)EtO	Me	H	5-Ind	G-107			
G-110	2(PhNH)EtO	H	H	5-Ind	G-108			
G-111	2(PhNH)EtO	Me	H	1Me-5-Ind	G-107			
G-112	2(PhNH)EtO	H	H	1Me-5-Ind	G-108	C		415(M <sup>+</sup> +1)
G-113	2(PhNH)EtO	Me	H	5-1HIdz	G-107			
G-114	2(PhNH)EtO	H	H	5-1HIdz	G-108			
G-115	2(PhNH)EtO	Me	H	1Me-5-1HIdz	G-107	A	4.76	430(M <sup>+</sup> +1)
G-116	2(PhNH)EtO	H	H	1Me-5-1HIdz	G-108	C		416(M <sup>+</sup> +1)
G-117	iBuO	H	H	1Me-5-Ind	G-1,G-2	C		352(M <sup>+</sup> +1)
G-118	iBuO	H	H	1Me-5-1HIdz	G-1,G-2	C		353(M <sup>+</sup> +1)
G-119	PhO	H	H	1Me-5-1HIdz	G-3,G-4	A	4.10	373(M <sup>+</sup> +1)
G-120	4ClPhO	H	H	1Me-5-1HIdz	G-3,G-4	A	4.46	407(M <sup>+</sup> +1)
G-121	4MeOPhO	H	H	1Me-5-1HIdz	G-3,G-4	A	4.12	403(M <sup>+</sup> +1)

[Examples H-1 to H-32]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification are shown in Table-H-1 and Table-H-2.

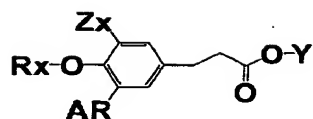
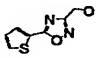
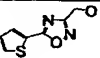
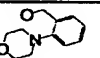
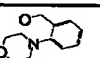
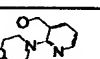
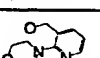
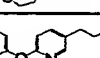
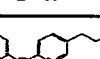
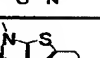
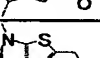
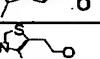
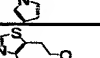
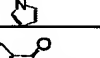
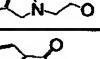


Table-H-1

Exp.	RxO	Y	Zx	AR	Syn	LCMS		
						method	RTime	Mass
H-1		Me	H	2-Nap	G-1			
H-2		H	H	2-Nap	G-2			
H-3		Me	H	5-Ind	G-1	C		375 (M <sup>+</sup> +1)
H-4		H	H	5-Ind	G-2			
H-5		Me	H	1Me-5-Ind	G-1			
H-6		H	H	1Me-5-Ind	G-2			
H-7		Me	H	5-1HIdz	G-1			
H-8		H	H	5-1HIdz	G-2			
H-9		Me	H	1Me-5-1HIdz	G-1			
H-10		H	H	1Me-5-1HIdz	G-2	C		454 (M <sup>+</sup> +1)
H-11		H	H	2-Nap	G-1,G-2			
H-12		H	H	1Me-5-Ind	G-1,G-2	C		452 (M <sup>+</sup> +1)
H-13		H	H	2-Nap	G-1,G-2			
H-14		H	H	1Me-5-Ind	G-1,G-2			
H-15		H	H	2-Nap	G-1,G-2	C		464 (M <sup>+</sup> +1)
H-16		H	H	1Me-5-Ind	G-1,G-2			
H-17		H	H	2-Nap	G-1,G-2	C		450 (M <sup>+</sup> +1)
H-18		H	H	1Me-5-Ind	G-1,G-2			



Table-H-2

H-19		H	H	2-Nap	G-1,G-2			
H-20		H	H	1Me-5-Ind	G-1,G-2			
H-21		H	H	2-Nap	G-1,G-2			
H-22		H	H	1Me-5-Ind	G-1,G-2	C		471 (M <sup>+</sup> +1)
H-23		H	H	2-Nap	G-1,G-2			
H-24		H	H	1Me-5-Ind	G-1,G-2			
H-25		H	H	2-Nap	G-1,G-2			
H-26		H	H	1Me-5-Ind	G-1,G-2			
H-27		H	H	2-Nap	G-1,G-2			
H-28		H	H	1Me-5-Ind	G-1,G-2	C		460 (M <sup>+</sup> +1)
H-29		H	H	2-Nap	G-1,G-2			
H-30		H	H	1Me-5-Ind	G-1,G-2			
H-31		H	H	2-Nap	G-1,G-2	C		452 (M <sup>+</sup> +1)
H-32		H	H	1Me-5-Ind	G-1,G-2			

## [Example J-1]

Synthesis of methyl 3-[4-cyclopentylmethoxy-3-fluoro-5-(1H-indol-5-yl)phenyl]propionate (Compound No. J-1)

According to the procedure described in the synthesis method of Compound No. C-1 with the modifications that the reaction was carried out for 13 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 10:1), Compound No. A-21 (154 mg), 5-indoleboronic acid (100 mg), 2 M aqueous sodium carbonate (1.5 ml) and (Ph<sub>3</sub>P)<sub>4</sub>Pd (50 mg) were reacted and treated to obtain the title compound (Compound No. J-1, 125 mg).

## [Example J-2]

Synthesis of 3-[4-cyclopentylmethyloxy-3-fluoro-5-(1H-indol-5-yl)phenyl]propionic acid (Compound No. J-2)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 2 hours, Compound No. J-1 (124 mg) and 2 N aqueous sodium hydroxide (630  $\mu$  l) were reacted and treated to obtain the title compound (Compound No. J-2, 97 mg).

[Example J-3]

Synthesis of methyl 3-[3-chloro-4-cyclopentylmethyloxy-5-(1H-indol-5-yl)phenyl]propionate (Compound No. J-3)

According to the procedure described in the synthesis method of Compound No. C-1 with the modifications that the reaction was carried out for 13 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 10:1), Compound No. A-20 (151 mg), 5-indoleboronic acid (97 mg), 2 M aqueous sodium carbonate (1.5 ml) and (Ph<sub>3</sub>P)<sub>4</sub>Pd (46 mg) were reacted and treated to obtain the title compound (Compound No. J-3, 160 mg).

[Example J-4]

Synthesis of 3-[3-chloro-4-cyclopentylmethyloxy-5-(1H-indol-5-yl)phenyl]propionic acid (Compound No. J-4)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 2 hours, Compound No. J-3 (135 mg) and 2 N aqueous sodium hydroxide (660  $\mu$  l) were reacted and treated to obtain the title compound (Compound No. J-4, 97 mg).

[Examples J-1 to J-92]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification including the examples described above are shown in Table-J-1 to Table-J-3

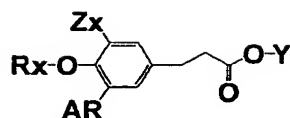


Table-J-1

Exp.	RxO	Y	Zx	AR	Syn	LCMS		
						method	RTime	Mass
J-1	cPenMeO	Me	F	5-Ind	J-1	A		396 (M <sup>+</sup> +1)
J-2	cPenMeO	H	F	5-Ind	J-2			
J-3	cPenMeO	Me	Cl	5-Ind	J-3			
J-4	cPenMeO	H	Cl	5-Ind	J-4	C		398 (M <sup>+</sup> +1)
J-5	cPenMeO	Me	F	2-Nap	J-1			
J-6	cPenMeO	H	F	2-Nap	J-2			
J-7	cPenMeO	Me	F	1Me-5-Ind	J-1			
J-8	cPenMeO	H	F	1Me-5-Ind	J-2			
J-9	cPenMeO	Me	F	5-1Hldz	J-1			
J-10	cPenMeO	H	F	5-1Hldz	J-2			
J-11	cPenMeO	Me	F	1Me-5-1Hldz	J-1			
J-12	cPenMeO	H	F	1Me-5-1Hldz	J-2	C		397 (M <sup>+</sup> +1)
J-13	2EtBuO	H	F	2-Nap	G-1,G-2			
J-14	2EtBuO	H	F	5-Ind	G-1,G-2			
J-15	4Me,cHexO	H	F	2-Nap	G-1,G-2			
J-16	4Me,cHexO	H	F	1Me-5-Ind	G-1,G-2			
J-17		H	F	2-Nap	G-1,G-2			
J-18		H	F	1Me-5-Ind	G-1,G-2	C		452 (M <sup>+</sup> +1)
J-19	cHepO	H	F	2-Nap	G-1,G-2			
J-20	3PhPrO	H	F	1Me-5-Ind	G-1,G-2			
J-21	4PhBuO	H	F	2-Nap	G-1,G-2			
J-22		H	F	1Me-5-Ind	G-1,G-2			
J-23	1(4MePh)EtO	H	F	2-Nap	G-1,G-2			
J-24	4ClBnO	H	F	1Me-5-Ind	G-1,G-2			
J-25	4CF3BnO	H	F	2-Nap	G-1,G-2			
J-26	3F,4(OMe)BnO	H	F	1Me-5-Ind	G-1,G-2			
J-27		H	F	2-Nap	G-1,G-2	C		429 (M <sup>+</sup> +1)
J-28		H	F	1Me-5-Ind	G-1,G-2			
J-29		H	F	1Me-5-Ind	G-1,G-2			
J-30		H	F	2-Nap	G-1,G-2			
J-31		H	F	2-Nap	G-1,G-2			

Table-J-2

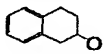

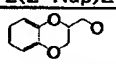
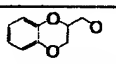
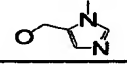
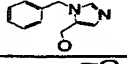
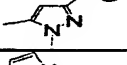
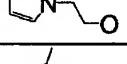
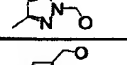
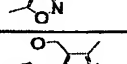
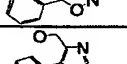
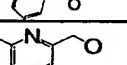
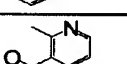
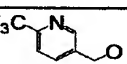
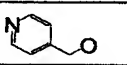
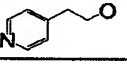
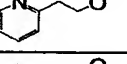
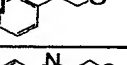
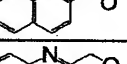
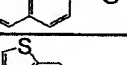
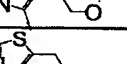
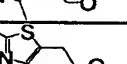
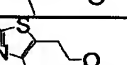

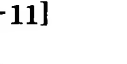
J-32	1-IndanO	H	F	2-Nap	G-1,G-2			
J-33	2-IndaneO	H	F	1Me-5-Ind	G-1,G-2			
J-34	2-IndaneO	H	F	2-Nap	G-1,G-2			
J-35	5OMe-2-IndanO	H	F	1Me-5-Ind	G-1,G-2			
J-36	5,6D(OMe)-2-IndanO	H	F	2-Nap	G-1,G-2			
J-37	5F-2-IndanO	H	F	2-Nap	G-1,G-2			
J-38	5F-2-IndanO	H	F	1Me-5-Ind	G-1,G-2			
J-39		H	F	2-Nap	G-1,G-2	C		441 (M <sup>+</sup> +1)
J-40		H	F	1Me-5-Ind	G-1,G-2			
J-41	2(3MePh)EtO	H	F	2-Nap	G-1,G-2			
J-42	2(4MePh)EtO	H	F	1Me-5-Ind	G-1,G-2			
J-43	2(2ClPh)EtO	H	F	1Me-5-Ind	G-1,G-2			
J-44	2(3ClPh)EtO	H	F	2-Nap	G-1,G-2			
J-45	2(2CF <sub>3</sub> Ph)EtO	H	F	2-Nap	G-1,G-2			
J-46	2(2OMePh)EtO	H	F	1Me-5-Ind	G-1,G-2			
J-47	2(4OMePh)EtO	H	F	2-Nap	G-1,G-2			
J-48	2(2-Nap)EtO	H	F	1Me-5-Ind	G-1,G-2			
J-49		H	F	2-Nap	G-1,G-2	C		458 (M <sup>+</sup> +1)
J-50		H	F	1Me-5-1HIdz	G-1,G-2			
J-51	2(PhS)EtO	H	F	1Me-5-Ind	G-1,G-2			
J-52	3PhPrO	H	F	2-Nap	G-1,G-2			
J-53	2ClBnO	H	F	1Me-5-Ind	G-1,G-2			
J-54	2BrBnO	H	F	2-Nap	G-1,G-2			
J-55	3,5DMeBnO	H	F	2-Nap	G-1,G-2			
J-56	4tBuBnO	H	F	1Me-5-Ind	G-1,G-2	C		460 (M <sup>+</sup> +1)
J-57	2CF <sub>3</sub> BnO	H	F	1Me-5-Ind	G-1,G-2			
J-58	4CF <sub>3</sub> BnO	H	F	2-Nap	G-1,G-2			
J-59	4nBuOBnO	H	F	2-Nap	G-1,G-2			
J-60	3,5DClBnO	H	F	1Me-5-Ind	G-1,G-2			
J-61	2,3DClBnO	H	F	2-Nap	G-1,G-2			
J-62	2-NapMeO	H	F	2-Nap	G-1,G-2	C		451 (M <sup>+</sup> +1)
J-63	1-NapMeO	H	F	1Me-5-Ind	G-1,G-2			
J-64	2PhBnO	H	F	1Me-5-Ind	G-1,G-2			
J-65	4PhBnO	H	F	1Me-5-Ind	G-1,G-2			
J-66	5OMe-2-IndanO	H	F	2-Nap	G-1,G-2			
J-67	5OMe-2-IndanO	H	F	1Me-5-Ind	G-1,G-2			
J-68	5,8D(OMe)-2-IndanO	H	F	2-Nap	G-1,G-2			
J-69	5,6D(OMe)-2-IndanO	H	F	1Me-5-Ind	G-1,G-2			
J-70	5F-2-IndanO	H	F	2-Nap	G-1,G-2			
J-71	5F-2-IndanO	H	F	1Me-5-Ind	G-1,G-2			

Table-J-3

J-72		H	F	1Me-5-Ind	G-1,G-2			
J-73		H	F	2-Nap	G-1,G-2	C		481 (M <sup>+</sup> +1)
J-74		H	F	1Me-5-Ind	G-1,G-2			
J-75		H	F	2-Nap	G-1,G-2			
J-76		H	F	1Me-5-Ind	G-1,G-2			
J-77		H	F	1Me-5-1HIdz	G-1,G-2	C		410 (M <sup>+</sup> +1)
J-78		H	F	1Me-5-Ind	G-1,G-2			
J-79		H	F	2-Nap	G-1,G-2			
J-80		H	F	1Me-5-Ind	G-1,G-2			
J-81		H	F	2-Nap	G-1,G-2			
J-82		H	F	1Me-5-Ind	G-1,G-2			
J-83		H	F	1Me-5-Ind	G-1,G-2			
J-84		H	F	1Me-5-Ind	G-1,G-2			
J-85		H	F	1Me-5-Ind	G-1,G-2	C		419 (M <sup>+</sup> +1)
J-86		H	F	1Me-5-Ind	G-1,G-2			
J-87		H	F	2-Nap	G-1,G-2			
J-88		H	F	1Me-5-Ind	G-1,G-2			
J-89		H	F	2-Nap	G-1,G-2	C		436 (M <sup>+</sup> +1)
J-90		H	F	1Me-5-Ind	G-1,G-2			
J-91		H	F	2-Nap	G-1,G-2			
J-92		H	F	1Me-5-Ind	G-1,G-2			

[Example K-11]

Synthesis of methyl 3-[3-bromo-4-cyclopentylmethoxy-5-(naphthalen-2-yl)phenyl]propionate (Compound No. K-11)

According to the procedure described in the synthesis method of Compound

No. C-1 with the modifications that the reaction was carried out for 15 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 9:1), Compound No. B-117 (306 mg), 2-naphthaleneboronic acid (163 mg), 2 M aqueous sodium carbonate (689  $\mu$ l) and  $(\text{Ph}_3\text{P})_4\text{Pd}$  (74.2 mg) were reacted and treated to obtain the title compound (Compound No. K-11, 261 mg).

Synthesis of 3-[3-bromo-4-cyclopentylmethoxy-5-(1H-indol-5-yl)phenyl]propionic acid (Compound No. K-12)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 2 hours, Compound No. K-11 (131 mg) and 2 N aqueous sodium hydroxide (400  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. K-12, 109 mg).

[Example K-13]

Synthesis of methyl 3-[3-bromo-4-cyclopentylmethoxy-5-(1H-indol-5-yl)phenyl]propionate (Compound No. K-13)

According to the procedure described in the synthesis method of Compound No. C-1 with the modifications that the reaction was carried out for 13 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 5:1), Compound No. B-117 (102 mg), 5-indoleboronic acid (97 mg), 2 M aqueous sodium carbonate (1.5 ml) and  $(\text{Ph}_3\text{P})_4\text{Pd}$  (46 mg) were reacted and treated to obtain the title compound (Compound No. K-13, 85 mg).

[Example K-14]

Synthesis of 3-[3-bromo-4-cyclopentylmethoxy-5-(1H-indol-5-yl)phenyl]propionic acid (Compound No. K-14)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 2 hours, Compound No. K-13 (85 mg) and 2 N aqueous sodium hydroxide (200  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. K-14, 79 mg).

[Example K-17]

Synthesis of methyl 3-[3-bromo-4-cyclopentyloxy-5-(1-methyl-1H-indazol-5-yl)phenyl]propionate (Compound No. K-17)

According to the procedure described in the synthesis method of Compound No. C-1 with the modifications that the reaction was carried out for 14 hours at 80°C, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 4:1), Compound No. B-118 (306 mg), 1-methyl-1H-indazole-5-boronic acid (175 mg), 2 M aqueous sodium carbonate (0.68 ml) and (Ph<sub>3</sub>P)<sub>4</sub>Pd (70.1 mg) were reacted and treated to obtain the title compound (Compound No. K-17, 148 mg).

[Examples K-1 to K-40]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification including the examples described above are shown in Table-K-1 and Table-K-2.

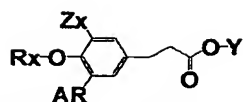
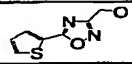
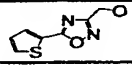
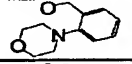
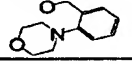
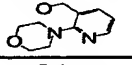
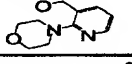
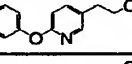
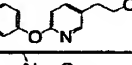
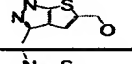
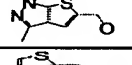


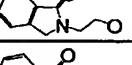



Table-K-1

Exp.	RxO	Y	Zx	AR	Syn	LCMS		
						method	RTime	Mass
K-1		Me	F	2-Nap	G-1			
K-2		H	F	2-Nap	G-2			
K-3		Me	F	5-Ind	G-1			
K-4		H	F	5-Ind	G-2	C		457(M <sup>+</sup> +1)
K-5		Me	F	1Me-5-Ind	G-1			
K-6		H	F	1Me-5-Ind	G-2	C		471(M <sup>+</sup> +1)
K-7		Me	F	5-1HIdz	G-1			
K-8		H	F	5-1HIdz	G-2			
K-9		Me	F	1Me-5-1HIdz	G-1			
K-10		H	F	1Me-5-1HIdz	G-2			
K-11	cPenMeO	Me	Br	2-Nap	K-11			
K-12	cPenMeO	H	Br	2-Nap	K-12			
K-13	cPenMeO	Me	Br	2-Nap	K-13			
K-14	cPenMeO	H	Br	5-Ind	Int50,K-13	C		456(M <sup>+</sup> )
K-15	cPenO	H	Br	2-Nap	K-11,K-12			
K-16	cPenO	H	Br	1Me-5-Ind	K-11,K-12			
K-17	cPenO	Me	Br	1Me-5-1HIdz	K-11,K-12			
K-18	cPenO	H	Br	1Me-5-1HIdz	K-11,K-12	A	4.78	443(M <sup>+</sup> )
K-19		H	F	2-Nap	G-1,G-2			
K-20		H	F	1Me-5-Ind	G-1,G-2			
K-21		H	F	2-Nap	G-1,G-2			
K-22		H	F	1Me-5-Ind	G-1,G-2			
K-23		H	F	2-Nap	G-1,G-2			
K-24		H	F	1Me-5-Ind	G-1,G-2	C		485(M <sup>+</sup> +1)
K-25		H	F	2-Nap	G-1,G-2			
K-26		H	F	1Me-5-Ind	G-1,G-2			



Table-K-2

K-27		H	F	2-Nap	G-1,G-2			
K-28		H	F	1Me-5-Ind	G-1,G-2			
K-29		H	F	2-Nap	G-1,G-2			
K-30		H	F	1Me-5-Ind	G-1,G-2			
K-31		H	F	2-Nap	G-1,G-2	C		486(M <sup>+</sup> +1)
K-32		H	F	1Me-5-Ind	G-1,G-2			
K-33		H	F	2-Nap	G-1,G-2			
K-34		H	F	1Me-5-Ind	G-1,G-2	C		511(M <sup>+</sup> +1)
K-35		H	F	2-Nap	G-1,G-2			
K-36		H	F	1Me-5-Ind	G-1,G-2			
K-37		H	F	2-Nap	G-1,G-2			
K-38		H	F	1Me-5-Ind	G-1,G-2			
K-39		H	F	2-Nap	G-1,G-2			
K-40		H	F	1Me-5-Ind	G-1,G-2			

## [Example L-1]

Synthesis of 3-[4-cyclopentyloxy-3-methyl-5-(naphthalen-2-yl)phenyl]propionic acid  
(Compound No. L-1)

According to the procedure described in the synthesis method of Compound No. C-1 with the modifications that the reaction was carried out at 80°C for 6 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 20:1), Compound A-24 (63 mg), 2-naphthaleneboronic acid (67 mg), 2 M aqueous sodium carbonate (130  $\mu$ l) and (Ph<sub>3</sub>P)<sub>4</sub>Pd (18 mg) were reacted and treated. The obtained substance was reacted with 2 N aqueous sodium hydroxide (200  $\mu$ l) and treated according to the procedure described in the synthesis method of Intermediate 9 to obtain the title compound (Compound No. L-1, 25 mg).

[Example L-2]

Synthesis of methyl 3-[4-cyclopentyloxy-3-methyl-5-(1-methyl-1H-indazol-5-yl)phenyl]propionate (Compound No. L-2)

According to the procedure described in the synthesis method of Compound No. C-1 with the modifications that the reaction was carried out at 80°C for 12 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 4:1), Compound No. K-17 (115 mg), methylboronic acid (66 mg, Ald), 2 M aqueous sodium carbonate (0.40 ml) and (Ph<sub>3</sub>P)<sub>4</sub>Pd (39.4 mg) were reacted and treated to obtain the title compound (Intermediate 52, 84 mg).

[Example L-3]

Synthesis of 3-[4-cyclopentyloxy-3-methyl-5-(1-methyl-1H-indazol-5-yl)phenyl]propionic acid (Compound No. L-3)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 1.5 hours, Compound No. L-2 (82 mg) and 2 N aqueous sodium hydroxide (0.26 ml) were reacted and treated to obtain the title compound (Compound No. L-3, 62 mg).

[Examples L-1 to L-95]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification including the examples described above are shown in Table-L-1 to Table-L-3.

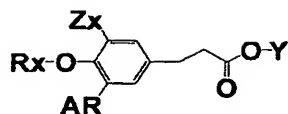


Table-L-1



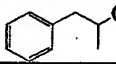
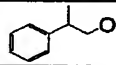
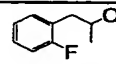
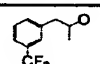
Exp.	RxO	Y	Zx	AR	Syn	LCMS		
						method	RTime	Mass
L-1	cPenO	H	Me	2-Nap	L-1	A	5.65	375(M <sup>+</sup> +1)
L-2	cPenO	Me	Me	1Me-5-1Hldz	L-2			
L-3	cPenO	H	Me	1Me-5-1Hldz	L-3	A	4.50	379(M <sup>+</sup> +1)
L-4	2EtBuO	Me	Me	2-Nap	L-2			
L-5	2EtBuO	H	Me	2-Nap	L-3	C		391(M <sup>+</sup> +1)
L-6	2EtBuO	H	Me	6-OMe-2-Nap	L-2,L-3			
L-7	2EtBuO	Me	Me	5-Ind	L-2			
L-8	2EtBuO	H	Me	5-Ind	L-3			
L-9	2EtBuO	Me	Me	1Me-5-Ind	L-2			
L-10	2EtBuO	H	Me	1Me-5-Ind	L-3			
L-11	2EtBuO	Me	Me	5-1Hldz	L-2			
L-12	2EtBuO	H	Me	5-1Hldz	L-3			
L-13	2EtBuO	Me	Me	1Me-5-1Hldz	L-2			
L-14	2EtBuO	H	Me	1Me-5-1Hldz	L-3	C		395(M <sup>+</sup> +1)
L-15	2EtBuO	Me	Me	5-Bzt	L-2			
L-16	2EtBuO	H	Me	5-Bzt	L-3			
L-17	2EtBuO	Me	Me	5-2ABzt	L-2			
L-18	2EtBuO	H	Me	5-2ABzt	L-3			
L-19	2EtBuO	Me	Me	2Me-5-Bzt	L-2			
L-20	2EtBuO	H	Me	2Me-5-Bzt	L-3			
L-21	4Me,cHexO	H	Me	1Me-5-Ind	G-1,G-2			
L-22		H	Me	2-Nap	G-1,G-2			
L-23	cHepO	H	Me	2-Nap	G-1,G-2			
L-24	cHepO	H	Me	1Me-5-Ind	G-1,G-2	C		406(M <sup>+</sup> +1)
L-25	3PhPrO	H	Me	2-Nap	G-1,G-2			
L-26	4PhBuO	H	Me	1Me-5-Ind	G-1,G-2			
L-27		H	Me	2-Nap	G-1,G-2			
L-28	1(4MePh)EtO	H	Me	1Me-5-Ind	G-1,G-2	C		428(M <sup>+</sup> +1)
L-29	4ClBnO	H	Me	2-Nap	G-1,G-2			
L-30	4CF <sub>3</sub> BnO	H	Me	1Me-5-Ind	G-1,G-2			
L-31	3F,4(OMe)BnO	H	Me	2-Nap	G-1,G-2			
L-32		H	Me	1Me-5-Ind	G-1,G-2			
L-33		H	Me	2-Nap	G-1,G-2			
L-34		H	Me	2-Nap	G-1,G-2			
L-35		H	Me	1Me-5-Ind	G-1,G-2			

Table-L-2

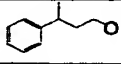
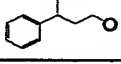
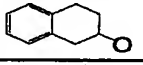

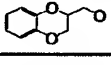
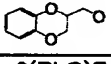
L-36		H	Me	1Me-5-Ind	G-1,G-2			
L-37		H	Me	1Me-5-Ind	G-1,G-2			
L-38	1-IndanO	H	Me	2-Nap	G-1,G-2			
L-39	2-IndanO	H	Me	2-Nap	G-1,G-2	C		423(M <sup>+</sup> +1)
L-40	2-IndanO	H	Me	1Me-5-Ind	G-1,G-2			
L-41	5OMe-2-IndanO	H	Me	1Me-5-Ind	G-1,G-2			
L-42	5,6D(OMe)-2-IndanO	H	Me	2-Nap	G-1,G-2			
L-43	5F-2-IndaneO	H	Me	2-Nap	G-1,G-2			
L-44	5F-2-IndaneO	H	Me	1Me-5-Ind	G-1,G-2			
L-45		H	Me	1Me-5-Ind	G-1,G-2			
L-46		H	Me	2-Nap	G-1,G-2			
L-47	2(3MePh)EtO	H	Me	2-Nap	G-1,G-2			
L-48	2(3FPh)EtO	H	Me	1Me-5-Ind	G-1,G-2	C		432(M <sup>+</sup> +1)
L-49	2(2ClPh)EtO	H	Me	1Me-5-Ind	G-1,G-2			
L-50	2(4CF <sub>3</sub> Ph)EtO	H	Me	1Me-5-Ind	G-1,G-2			
L-51	2(2OMePh)EtO	H	Me	2-Nap	G-1,G-2	C		441(M <sup>+</sup> +1)
L-52	2(4OMePh)EtO	H	Me	1Me-5-Ind	G-1,G-2			
L-53	2(2-Nap)EtO	H	Me	2-Nap	G-1,G-2			
L-54	2(2-Nap)EtO	H	Me	1Me-5-Ind	G-1,G-2			
L-55		H	Me	1Me-5-Ind	G-1,G-2			
L-56		H	Me	1Me-5-1HIdz	G-1,G-2	C		459(M <sup>+</sup> +1)
L-57	2(PhS)EtO	H	Me	2-Nap	G-1,G-2			
L-58	3PhPrO	H	Me	1Me-5-Ind	G-1,G-2			
L-59	2ClBnO	H	Me	2-Nap	G-1,G-2			
L-60	2BrBnO	H	Me	1Me-5-Ind	G-1,G-2			
L-61	3,5DMeBnO	H	Me	2-Nap	G-1,G-2			
L-62	4tBuBnO	H	Me	2-Nap	G-1,G-2			
L-63	2CF <sub>3</sub> BnO	H	Me	2-Nap	G-1,G-2			
L-64	4tBuBnO	H	Me	1Me-5-Ind	G-1,G-2			
L-65	4nBuBnO	H	Me	2-Nap	G-1,G-2	C		453(M <sup>+</sup> +1)
L-66	3,5DClBnO	H	Me	2-Nap	G-1,G-2			
L-67	2,3DClBnO	H	Me	1Me-5-Ind	G-1,G-2			
L-68	2-NapMeO	H	Me	1Me-5-Ind	G-1,G-2			
L-69	1-NapMeO	H	Me	2-Nap	G-1,G-2			
L-70	2PhBnO	H	Me	1Me-5-Ind	G-1,G-2			
L-71	4PhBnO	H	Me	2-Nap	G-1,G-2	C		476(M <sup>+</sup> +1)
L-72	5OMe-2-IndanO	H	Me	1Me-5-Ind	G-1,G-2			
L-73	5F-2-IndaneO	H	Me	2-Nap	G-1,G-2			

Table-L-3

L-74		H	Me	2-Nap	G-1,G-2	C		401(M <sup>+</sup> +1)
L-75		H	Me	1Me-5-Ind	G-1,G-2			
L-76		H	Me	2-Nap	G-1,G-2			
L-77		H	Me	1Me-5-Ind	G-1,G-2			
L-78		H	Me	2-Nap	G-1,G-2			
L-79		H	Me	1Me-5-Ind	G-1,G-2			
L-80		H	Me	2-Nap	G-1,G-2			
L-81		H	Me	1Me-5-Ind	G-1,G-2			
L-82		H	Me	2-Nap	G-1,G-2	C		412(M <sup>+</sup> +1)
L-83		H	Me	1Me-5-Ind	G-1,G-2			
L-84		H	Me	2-Nap	G-1,G-2			
L-85		H	Me	1Me-5-Ind	G-1,G-2			
L-86		H	Me	2-Nap	G-1,G-2			
L-87		H	Me	1Me-5-Ind	G-1,G-2			
L-88		H	Me	2-Nap	G-1,G-2			
L-89		H	Me	1Me-5-Ind	G-1,G-2	C		415(M <sup>+</sup> +1)
L-90		H	Me	2-Nap	G-1,G-2			
L-91		H	Me	1Me-5-Ind	G-1,G-2			
L-92		H	Me	2-Nap	G-1,G-2			
L-93		H	Me	1Me-5-Ind	G-1,G-2	C		435(M <sup>+</sup> +1)
L-94		H	Me	2-Nap	G-1,G-2			
L-95		H	Me	1Me-5-Ind	G-1,G-2			

[Examples M-1 to M-32]

Typical examples of the compounds of the present invention that can be

obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification are shown in Table-M-1 and Table-M-2.

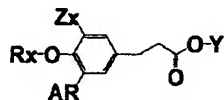
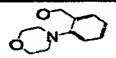

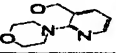
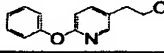
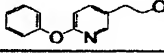
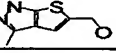



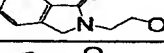
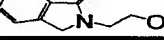


Table-M-1

Exp.	RxO	Y	Zx	AR	Syn	LCMS		
						method	RTime	Mass
M-1		Me	Me	2-Nap	G-1	C		478(M <sup>+</sup> +1)
M-2		H	Me	2-Nap	G-2	C		464(M <sup>+</sup> +1)
M-3		Me	Me	5-Ind	G-1			
M-4		H	Me	5-Ind	G-2			
M-5		Me	Me	1Me-5-Ind	G-1			
M-6		H	Me	1Me-5-Ind	G-2	C		467(M <sup>+</sup> +1)
M-7		Me	Me	5-1HIdz	G-1			
M-8		H	Me	5-1HIdz	G-2			
M-9		Me	Me	1Me-5-1HIdz	G-1			
M-10		H	Me	1Me-5-1HIdz	G-2			
M-11		H	Me	2-Nap	G-1,G-2	C		463(M <sup>+</sup> +1)
M-12		H	Me	1Me-5-Ind	G-1,G-2			
M-13		H	Me	2-Nap	G-1,G-2			
M-14		H	Me	1Me-5-Ind	G-1,G-2	C		465(M <sup>+</sup> +1)
M-15		H	Me	2-Nap	G-1,G-2			
M-16		H	Me	1Me-5-Ind	G-1,G-2			
M-17		H	Me	2-Nap	G-1,G-2	C		464(M <sup>+</sup> +1)
M-18		H	Me	1Me-5-Ind	G-1,G-2			
M-19		H	Me	2-Nap	G-1,G-2			
M-20		H	Me	1Me-5-Ind	G-1,G-2			
M-21		H	Me	2-Nap	G-1,G-2			

Table-M-2

M-22		H	Me	1Me-5-Ind	G-1,G-2			
M-23		H	Me	2-Nap	G-1,G-2			
M-24		H	Me	1Me-5-Ind	G-1,G-2	C		486(M <sup>+</sup> +1)
M-25		H	Me	2-Nap	G-1,G-2			
M-26		H	Me	1Me-5-Ind	G-1,G-2			
M-27		H	Me	2-Nap	G-1,G-2			
M-28		H	Me	1Me-5-Ind	G-1,G-2			
M-29		H	Me	2-Nap	G-1,G-2			
M-30		H	Me	1Me-5-Ind	G-1,G-2	C		472(M <sup>+</sup> +1)
M-31		H	Me	2-Nap	G-1,G-2			
M-32		H	Me	1Me-5-Ind	G-1,G-2			

## [Example N-1]

Synthesis of methyl 3-{4-[2-(N-acetyl-N-phenylamino)ethoxy]-3-(naphthalen-2-yl)phenyl}propionate (Compound No. N-1)

A solution of Compound No. G-107 (32 mg) in methylene chloride (1 ml) was added with pyridine (24  $\mu$ l, TCI) and acetyl chloride (21  $\mu$ l, TCI), and stirred for 17 hours. The reaction mixture was added with water (3 ml), and extracted with methylene chloride (10 ml). The organic layer was washed with saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 2:1) to obtain the title compound (Compound No. N-1, 28.1 mg).

## [Example N-2]

Synthesis of 3-{4-[2-(N-acetyl-N-phenylamino)ethoxy]-3-(naphthalen-2-yl)phenyl}propionic acid (Compound No. N-2)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 3 hours, Compound

No. N-1 (28 mg) and 2 N aqueous sodium hydroxide (0.25 ml) were reacted and treated to obtain the title compound (Compound No. N-2, 22 mg).

[Example N-29]

Synthesis of methyl 3-{4-[2-(N-methoxycarbonyl-N-phenylamino)ethyloxy]-3-(naphthalen-2-yl)phenyl}propionate (Compound No. N-29)

According to the procedure described in the synthesis method of Compound No. N-1, Compound No. G-107 (32 mg), pyridine (23  $\mu$ l) and methyl chloroformate (23  $\mu$ l, TCI) were reacted and treated to obtain the title compound (Compound No. N-29, 17.3 mg).

[Example N-30]

Synthesis of 3-{4-[2-(N-methoxycarbonyl-N-phenylamino)ethyloxy]-3-(naphthalen-2-yl)phenyl}propionic acid (Compound No. N-30)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 3 hours, Compound No. N-29 (17 mg) and 2 N aqueous sodium hydroxide (0.25 ml) were reacted and treated to obtain the title compound (Compound No. N-30, 10.1 mg).

[Example N-48]

Synthesis of methyl 3-{4-[2-(N-methylsulfonyl-N-phenylamino)ethyloxy]-3-(naphthalen-2-yl)phenyl}propionate (Compound No. N-48)

According to the procedure described in the synthesis method of Compound No. N-1, Compound No. G-107 (32 mg), pyridine (24  $\mu$ l) and methanesulfonyl chloride (23  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. N-48, 32.3 mg).

[Example N-49]

Synthesis of 3-{4-[2-(N-methylsulfonyl-N-phenylamino)ethyloxy]-3-(naphthalen-2-yl)phenyl}propionic acid (Compound No. N-49)

According to the procedure described in the synthesis method of



Intermediate 9 provided that the reaction was carried out for 3 hours, Compound No. N-48 (32 mg) and 2 N aqueous sodium hydroxide (0.25 ml) were reacted and treated to obtain the title compound (Compound No. N-49, 17 mg).

[Example N-55]

Synthesis of methyl 3-{4-[2-(3-ethyl-1-phenylureido)ethyloxy]-3-(naphthalen-2-yl)phenyl}propionate (Compound No. N-55)

According to the procedure described in the synthesis method of Compound No. N-1 provided that the reaction was carried out for 41 hours, Compound No. G-107 (32 mg), pyridine (24  $\mu$ l) and ethyl isocyanate (24  $\mu$ l, Nakarai Tecs) were reacted and treated to obtain the title compound (Compound No. N-55, 31.2 mg).

[Example N-56]

Synthesis of 3-{4-[2-(3-ethyl-1-phenylureido)ethyloxy]-3-(naphthalen-2-yl)phenyl}propionic acid (Compound No. N-56)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 3 hours, Compound No. N-55 (31 mg) and 2 N aqueous sodium hydroxide (0.25 ml) were reacted and treated to obtain the title compound (Compound No. N-56, 15 mg).

[Example N-64]

Synthesis of methyl 3-{4-[2-(3-ethyl-1-phenylthioureido)ethyloxy]-3-(naphthalen-2-yl)phenyl}propionate (Compound No. N-64)

According to the procedure described in the synthesis method of Compound No. N-1 provided that the reaction was carried out for 41 hours, Compound No. G-107 (32 mg), pyridine (24  $\mu$ l) and ethyl isothiocyanate (21  $\mu$ l, Nakarai Tecs) were reacted and treated to obtain the title compound (Compound No. N-64, 27.4 mg).

[Example N-65]

Synthesis of 3-{4-[2-(3-ethyl-1-phenylthioureido)ethyloxy]-3-(naphthalen-2-yl)phenyl}propionic acid (Compound No. N-65)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 3 hours, Compound No. N-64 (27 mg) and 2 N aqueous sodium hydroxide (0.25 ml) were reacted and treated to obtain the title compound (Compound No. N-65, 8.9 mg).

[Examples N-1 to N-74]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification including the examples described above are shown in Table-N-1 and Table-N-2.

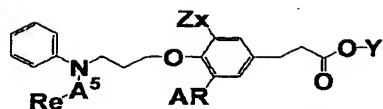


Table-N-1

Exp.	A <sup>5</sup> Re	Y	Zx	AR	Syn	LCMS		
						method	RTime	Mass
N-1	COMe	Me	H	2-Nap	N-1			
N-2	COMe	H	H	2-Nap	N-2			
N-3	COMe	Me	H	5-Ind	N-1			
N-4	COMe	H	H	5-Ind	N-2	C		457(M <sup>+</sup> +1)
N-5	COMe	Me	H	1Me-5-Ind	N-1			
N-6	COMe	H	H	1Me-5-Ind	N-2			
N-7	COMe	Me	H	5-1HIdz	N-1			
N-8	COMe	H	H	5-1HIdz	N-2			
N-9	COMe	Me	H	1Me-5-1HIdz	N-1			
N-10	COMe	H	H	1Me-5-1HIdz	N-2			
N-11	COPh	H	H	2-Nap	N-1,N-2	C		516(M <sup>+</sup> +1)
N-12	COPh	H	H	1Me-5-Ind	N-1,N-2			
N-13	COTBu	H	H	2-Nap	N-1,N-2			
N-14	COTBu	H	H	1Me-5-Ind	N-1,N-2			
N-15	COiPr	H	H	2-Nap	N-1,N-2	C		496(M <sup>+</sup> +1)
N-16	COiPr	H	H	1Me-5-Ind	N-1,N-2			
N-17	COCH(Et) <sub>n</sub> Bu	H	H	2-Nap	N-1,N-2			
N-18	COCH(Et) <sub>n</sub> Bu	H	H	1Me-5-Ind	N-1,N-2			
N-19	COCH <sub>2</sub> OMe	H	H	2-Nap	N-1,N-2			
N-20	COCH <sub>2</sub> OMe	H	H	1Me-5-Ind	N-1,N-2			
N-21	COCH=CHMe	H	H	2-Nap	N-1,N-2			
N-22	COCH=CHMe	H	H	1Me-5-Ind	N-1,N-2	C		483(M <sup>+</sup> +1)
N-23	COiBu	H	H	2-Nap	N-1,N-2			
N-24	COiBu	H	H	1Me-5-Ind	N-1,N-2			
N-25	COcPr	H	H	2-Nap	N-1,N-2			
N-26	COcPr	H	H	1Me-5-Ind	N-1,N-2	C		483(M <sup>+</sup> +1)
N-27	CO(CH <sub>2</sub> ) <sub>2</sub> cPen	H	H	2-Nap	N-1,N-2			
N-28	CO(CH <sub>2</sub> ) <sub>2</sub> cPen	H	H	1Me-5-Ind	N-1,N-2			
N-29	COOMe	Me	H	2-Nap	N-29			
N-30	COOMe	H	H	2-Nap	N-30			
N-31	COOMe	H	H	1Me-5-Ind	N-29,N-30			
N-32	COOPh	H	H	2-Nap	N-29,N-30	C		516(M <sup>+</sup> +1)
N-33	COOPh	H	H	1Me-5-Ind	N-29,N-30			
N-34	CONMe <sub>2</sub>	H	H	2-Nap	N-29,N-30	C		483(M <sup>+</sup> +1)
N-35	CONMe <sub>2</sub>	H	H	1Me-5-Ind	N-29,N-30			
N-36	COOiBu	H	H	2-Nap	N-29,N-30			
N-37	COOiBu	H	H	1Me-5-Ind	N-29,N-30			
N-38	C(O)SMe	H	H	2-Nap	N-29,N-30			
N-39	C(O)SMe	H	H	1Me-5-Ind	N-29,N-30			
N-40		H	H	2-Nap	N-29,N-30			
N-41		H	H	1Me-5-Ind	N-29,N-30	C		528(M <sup>+</sup> +1)

Table-N-2

N-42		H	H	2-Nap	Int53,N-29			
N-43		H	H	1Me-5-Ind	Int53,N-29			
N-44	COO(CH <sub>2</sub> ) <sub>2</sub> OMe	H	H	2-Nap	Int53,N-29			
N-45	COO(CH <sub>2</sub> ) <sub>2</sub> OMe	H	H	1Me-5-Ind	Int53,N-29			
N-46		H	H	2-Nap	Int53,N-29			
N-47		H	H	1Me-5-Ind	Int53,N-29			
N-48	SO <sub>2</sub> Me	Me	H	2-Nap	N-48			
N-49	SO <sub>2</sub> Me	H	H	2-Nap	N-49			
N-50	SO <sub>2</sub> Me	H	H	1Me-5-Ind	N-48,N-49	C		493(M <sup>+</sup> +1)
N-51	SO <sub>2</sub> Ph	H	H	2-Nap	N-48,N-49			
N-52	SO <sub>2</sub> Ph	H	H	1Me-5-Ind	N-48,N-49			
N-53	SO <sub>2</sub> NMe <sub>2</sub>	H	H	2-Nap	N-48,N-49	C		519(M <sup>+</sup> +1)
N-54	SO <sub>2</sub> NMe <sub>2</sub>	H	H	1Me-5-Ind	N-48,N-49			
N-55	CONHEt	Me	H	2-Nap	N-55			
N-56	CONHEt	H	H	2-Nap	N-56	C		483(M <sup>+</sup> +1)
N-57	CONHEt	H	H	1Me-5-Ind	N-55,N-56			
N-58	CONHPh	H	H	2-Nap	N-55,N-56			
N-59	CONHPh	H	H	1Me-5-Ind	N-55,N-56			
N-60	CONHcHex	H	H	2-Nap	N-55,N-56			
N-61	CONHcHex	H	H	1Me-5-Ind	N-55,N-56	C		540(M <sup>+</sup> +1)
N-62	CONHBn	H	H	2-Nap	N-55,N-56			
N-63	CONHBn	H	H	1Me-5-Ind	N-55,N-56			
N-64	CSNHMe	Me	H	2-Nap	N-64			
N-65	CSNHMe	H	H	2-Nap	N-65			
N-66	CSNHMe	H	H	1Me-5-Ind	N-64,N-65			
N-67	CSNHPh	H	H	2-Nap	N-64,N-65			
N-68	CSNHPh	H	H	1Me-5-Ind	N-64,N-65			
N-69	CSNH(3-Py)	H	H	2-Nap	N-64,N-65	C		548(M <sup>+</sup> +1)
N-70	CSNH(3-Py)	H	H	1Me-5-Ind	N-64,N-65			
N-71	CSNH <i>i</i> Pr	H	H	2-Nap	N-64,N-65			
N-72	CSNH <i>i</i> Pr	H	H	1Me-5-Ind	N-64,N-65	C		516(M <sup>+</sup> +1)
N-73	CSNHBn	H	H	2-Nap	N-64,N-65			
N-74	CSNHBn	H	H	1Me-5-Ind	N-64,N-65			

## [Example P-1]

Synthesis of ethyl 3-[2-cyclopentylmethoxy-3-(naphthalen-2-yl)pyridin-5-yl]propionate (Compound No. P-1)

According to the procedure described in the synthesis method of Compound No. C-1 with the modifications that the reaction was carried out for 14 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 5:1), 2-naphthaleneboronic acid (119 mg), Compound No. E-1 (83 mg), 2 M

aqueous sodium carbonate (0.3 ml) and  $(\text{Ph}_3\text{P})_4\text{Pd}$  (38.1 mg) were reacted and treated to obtain the title compound (Compound No. P-1, 76 mg).

[Example P-2]

Synthesis of 3-[2-cyclopentylmethoxy-3-(naphthalen-2-yl)pyridin-5-yl]propionic acid (Compound No. P-2)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 2 hours, Compound No. P-1 (47.8 mg) and 2 N aqueous sodium hydroxide (0.2 ml) were reacted and treated to obtain the title compound (Compound No. P-2, 20 mg).

[Example P-36]

Synthesis of ethyl 3-{3-(naphthalen-2-yl)-2-[(R)-1-phenylethyloxy]pyridin-5-yl}propionate (Compound No. P-36)

According to the procedure described in the synthesis method of Compound No. C-1 with the modifications that the reaction was carried out for 2 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 6:1), 2-naphthaleneboronic acid (44 mg), Compound No. E-7 (73.3 mg), 2 M aqueous sodium carbonate (120  $\mu$ l) and  $(\text{Ph}_3\text{P})_4\text{Pd}$  (21.3 mg) were reacted and treated to obtain the title compound (Compound No. P-36, 44 mg).

[Example P-37]

Synthesis of 3-{3-(naphthalen-2-yl)-2-[(R)-1-phenylethyloxy]pyridin-5-yl}propionic acid (Compound No. P-37)

According to the procedure described in the synthesis method of Intermediate 9, Compound No. P-36 (41.2 mg) and 2 N aqueous sodium hydroxide (0.1 ml) were reacted and treated to obtain the title compound (Compound No. P-37, 38 mg).

[Example P-42]

Synthesis of ethyl 3-{3-(naphthalen-2-yl)-2-[4-

(trifluoromethyl)phenylmethyloxy]pyridin-5-yl}propionate (Compound No. P-42)

According to the procedure described in the synthesis method of Compound No. C-1 with the modifications that the reaction was carried out for 2 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 6:1), 2-naphthaleneboronic acid (37.4 mg), Compound No. E-13 (42.4 mg), 2 M aqueous sodium carbonate (90  $\mu$ l) and  $(\text{Ph}_3\text{P})_4\text{Pd}$  (21.4 mg) were reacted and treated to obtain the title compound (Compound No. P-42, 30.4 mg).

[Example P-43]

Synthesis of 3-{3-(naphthalen-2-yl)-2-[4-(trifluoromethyl)phenylmethyloxy]pyridin-5-yl}propionic acid (Compound No. P-43)

According to the procedure described in the synthesis method of Intermediate 9, Compound No. P-42 (29.5 mg) and 2 N aqueous sodium hydroxide (0.15 ml) were reacted and treated to obtain the title compound (Compound No. P-43, 24.1 mg).

[Examples P-1 to P-50]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification including the examples described above are shown in Table-P-1 and Table-P-2.

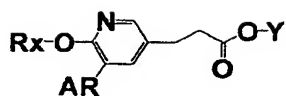


Table-P-1

Exp.	RxO	Y	AR	Syn	LCMS		
					method	RTime	Mass
P-1	cPenMeO	Et	2-Nap	P-1			
P-2	cPenMeO	H	2-Nap	P-2	A	5.60	376(M <sup>+</sup> +1)
P-3	cPenMeO	Et	5-Ind	P-1	A	5.37	393(M <sup>+</sup> +1)
P-4	cPenMeO	H	5-Ind	P-2			
P-5	cPenMeO	Et	1Me-5-Ind	P-1			
P-6	cPenMeO	H	1Me-5-Ind	P-2	A	4.90	379(M <sup>+</sup> +1)
P-7	cPenMeO	Et	1Me-5-Ind	P-1			
P-8	cPenMeO	H	5-1HIdz	P-2			
P-9	cPenMeO	Et	5-1HIdz	P-1			
P-10	cPenMeO	H	1Me-5-1HIdz	P-2			
P-11	cPenMeO	Et	5-Bzt	P-1			
P-12	cPenMeO	H	5-Bzt	P-2			
P-13	cPenMeO	Et	5-2ABzt	P-1			
P-14	cPenMeO	H	5-2ABzt	P-2			
P-15	cPenMeO	H	6-IQ	P-1,P-2	C		377(M <sup>+</sup> +1)
P-16	cPenO	H	2-Nap	P-1,P-2			
P-17	cPenO	H	5-Ind	P-1,P-2	C		351(M <sup>+</sup> +1)
P-18	cPenO	H	1Me-5-Ind	P-1,P-2			
P-19	cPenO	H	5-1HIdz	P-1,P-2			
P-20	cPenO	H	1Me-5-1HIdz	P-1,P-2			
P-21	cPenO	H	5-Bzt	P-1,P-2			
P-22	cPenO	H	5-2ABzt	P-1,P-2			
P-23	cHexO	H	2-Nap	P-1,P-2	A	5.51	376(M <sup>+</sup> +1)
P-24	cHexO	H	5-Ind	P-1,P-2			
P-25	cHexO	H	1Me-5-Ind	P-1,P-2			
P-26	cHexO	H	1Me-5-1HIdz	P-1,P-2			
P-27	2EtBuO	H	2-Nap	P-1,P-2	A	5.68	378(M <sup>+</sup> +1)
P-28	2EtBuO	H	5-Ind	P-1,P-2			
P-29	2EtBuO	H	1Me-5-Ind	P-1,P-2			
P-30	iBuO	H	2-Nap	P-1,P-2	A	5.13	350(M <sup>+</sup> +1)
P-31	iBuO	H	5-Ind	P-1,P-2			
P-32	iBuO	H	1Me-5-Ind	P-1,P-2			
P-33	iBuO	H	1Me-5-1HIdz	P-1,P-2			
P-34	BnO	H	2-Nap	P-1,P-2			
P-35	BnO	H	1Me-5-Ind	P-1,P-2			
P-36	(R)1PhEtO	Et	2-Nap	P-36			
P-37	(R)1PhEtO	H	2-Nap	P-37			
P-38	(S)1PhEtO	H	2-Nap	P-36,P37	A	5.31	398(M <sup>+</sup> +1)
P-39	(S)1PhEtO	H	1Me-5-Ind	P-36,P37	A	4.75	401(M <sup>+</sup> +1)
P-40	2MeBnO	H	2-Nap	P-1,P-2			
P-41	2MeBnO	H	1Me-5-Ind	P-1,P-2			

Table-P-2

P-42	4CF3BnO	Et	2-Nap	P-42			
P-43	4CF3BnO	H	2-Nap	P-43	A	5.52	452(M <sup>+</sup> +1)
P-44	4CF3BnO	H	1Me-5-Ind	P-1,P-2			
P-45	3PhBuO	H	1Me-5-Ind	P-1,P-2			
P-46	2(2-Nap)EtO	H	2-Nap	P-1,P-2			
P-47	2(2-Nap)EtO	H	1Me-5-Ind	P-1,P-2			
P-48	2(2FPh)EtO	H	2-Nap	P-1,P-2			
P-49	2(2FPh)EtO	H	5-Ind	P-1,P-2	A	4.18	405(M <sup>+</sup> +1)
P-50	2(2FPh)EtO	H	1Me-5-Ind	P-1,P-2			

[Example Q-1]

Synthesis of methyl 3-[4-methoxy-3-(naphthalen-2-yl)-5-nitrophenyl]propionate  
(Intermediate 49)

According to the procedure described in the synthesis method of Compound No. C-1 with the modifications that the reaction was carried out at 80°C for 15 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 10:1), Intermediate 21 (2.65 g), 2-naphthaleneboronic acid (2.87 g), 2 M aqueous sodium carbonate (7.5 ml) and (Ph<sub>3</sub>P)<sub>4</sub>Pd (960 mg) were reacted and treated to obtain the title compound (Intermediate 49, 2.47 g).  
Synthesis of 3-[4-methoxy-3-(naphthalen-2-yl)-5-nitrophenyl]propionic acid  
(Intermediate 50)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 40 minutes, Intermediate 49 (2.45 g) and 2 N aqueous sodium hydroxide (6.7 ml) were reacted and treated to obtain the title compound (Intermediate 60, 1.96 g).  
Synthesis of methyl 3-[4-hydroxy-3-(naphthalen-2-yl)-5-nitrophenyl]propionate  
(Intermediate 51)

According to the procedure described in the synthesis method of Intermediate 10 provided that the reaction was carried out for 3 hours, pyridine (10 ml), concentrated hydrochloric acid (10 ml), and Intermediate 50 (1.00 g) were reacted and treated to obtain crude powder substance. This substance was reacted



with thionyl chloride (282  $\mu$ l) in methanol and treated according to the procedure described in the synthesis method of Intermediate 1 to obtain the title compound (Intermediate 51, 306 mg).

Synthesis of methyl 3-[4-cyclopentyloxy-3-(naphthalen-2-yl)-5-nitrophenyl]propionate (Compound No. Q-1)

According to the procedure described in the synthesis method of Compound No. A-6 with the modifications that the reaction was carried out for 15.5 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 19:1), Intermediate 51 (84 mg),  $\text{Ph}_3\text{P}$  (125 mg), cyclopentanol (50  $\mu$ l) and 40% DIAD (224  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. Q-1, 90 mg).

[Example Q-2]

Synthesis of methyl 3-[3-amino-4-cyclopentyloxy-5-(naphthalen-2-yl)phenyl]propionate (Compound No. Q-2)

A solution of Compound No. Q-1 (59.1 mg) in methanol (5 ml) was added with platinum oxide (5 mg, Ald), and stirred at room temperature for 30 minutes under hydrogen atmosphere. The reaction mixture was filtered, and the solvent of the filtrate was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 4:1) to obtain the title compound (Compound No. Q-2, 49 mg).

[Example Q-3]

Synthesis of 3-[3-amino-4-cyclopentyloxy-5-(naphthalen-2-yl)phenyl]propionic acid (Compound No. Q-3)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 2 hours, Compound No. Q-2 (40 mg) and 2 N aqueous sodium hydroxide (150  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. Q-3, 38 mg).

[Example Q-4]

Synthesis of methyl 3-[4-cyclopentyloxy-3-(1H-indol-5-yl)-5-nitrophenyl]propionate  
(Compound No. Q-4)

According to the procedure described in the synthesis method of Compound No. C-1 with the modifications that the reaction was carried out at 80°C for 16 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 4:1), Compound No. A-28 (187 mg), 5-indoleboronic acid (143 mg), 2 M aqueous sodium carbonate (400  $\mu$ l) and (Ph<sub>3</sub>P)<sub>4</sub>Pd (51 mg) were reacted and treated to obtain the title compound (Compound No. Q-4, 192 mg).

[Example Q-5]

Synthesis of methyl 3-[3-amino-4-cyclopentyloxy-5-(1H-indol-5-yl)phenyl]propionate  
(Compound No. Q-5)

According to the procedure described in the synthesis method of Compound No. Q-2 with the modification that the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 2:1), Compound No. Q-4 (59.1 mg) and platinum oxide (5 mg) were reacted and treated to obtain the title compound (Compound No. Q-5, 49.3 mg).

[Example Q-6]

Synthesis of 3-[3-amino-4-cyclopentyloxy-5-(1H-indol-5-yl)phenyl]propionic acid  
(Compound No. Q-6)

According to the procedure described in the synthesis method of Intermediate 9, Compound No. Q-5 (44 mg) and 2 N aqueous sodium hydroxide (150  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. Q-6, 41 mg).

[Example Q-8]

Synthesis of methyl 3-[4-cyclopentyloxy-3-(1-methyl-1H-indazol-5-yl)-5-nitrophenyl]propionate (Compound No. Q-8)

According to the procedure described in the synthesis method of Compound No. C-1 with the modifications that the reaction was carried out at 80°C for 16 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 3:1), Compound No. A-28 (182 mg), 1-methyl-5-indazoleboronic acid (152 mg), 2 M aqueous sodium carbonate (400  $\mu$ l) and (Ph<sub>3</sub>P)<sub>4</sub>Pd (58.9 mg) were reacted and treated to obtain the title compound (Compound No. Q-8, 181 mg).

[Example Q-9]

Synthesis of methyl 3-[3-amino-4-cyclopentyloxy-5-(1-methyl-1H-indazol-5-yl)phenyl]propionic acid (Compound No. Q-9)

A solution of Compound No. Q-8 (578 mg) in a mixture of ethyl acetate (2 ml) and methanol (5 ml) was added with Raney 2800 nickel (230 mg) and stirred at room temperature for 6 hours under hydrogen atmosphere. The reaction mixture was filtered, and the solvent of the filtrate was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 2:1) to obtain the title compound (Compound No. Q-9, 484 mg).

[Example Q-10]

Synthesis of 3-[3-amino-4-cyclopentyloxy-5-(1H-indazol-5-yl)phenyl]propionic acid (Compound No. Q-10)

According to the procedure described in the synthesis method of Intermediate 9, Compound No. Q-9 (56 mg) and 2 N aqueous sodium hydroxide (200  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. Q-10, 50 mg).

[Example Q-47]

Synthesis of methyl 3-[4-benzyloxy-3-(naphthalen-2-yl)-5-nitrophenyl]propionate (Compound No. Q-47)

According to the procedure described in the synthesis method of Compound

No. C-1 with the modifications that the reaction was carried out at 80°C for 12 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 8:1), Compound No. B-95 (6.00 g), 2-naphthaleneboronic acid (4.11 g), 2 M aqueous sodium carbonate (13.5 ml) and (Ph<sub>3</sub>P)<sub>4</sub>Pd (1.36 g) were reacted and treated to obtain the title compound (Compound No. Q-47, 5.81 g).  
[Example Q-48]

Synthesis of methyl 3-[3-amino-4-benzyloxy-5-(naphthalen-2-yl)phenyl]propionate (Compound No. Q-48)

According to the procedure described in the synthesis method of Compound No. Q-9 with the modifications that the reaction was carried out for 20 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 2:1), Compound No. Q-47 (5.04 g) and Raney 2800 nickel (2.50 g) were reacted and treated to obtain the title compound (Compound No. Q-48, 4.21 g).  
[Example Q-1 to Q-52]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification including the examples described above are shown in Table-Q-1.

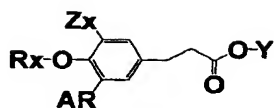


Table-Q-1

Exp.	RxO	Y	Zx	AR	Syn	LCMS		
						method	RTime	Mass
Q-1	cPenO	Me	NO2	2-Nap	Q-1			
Q-2	cPenO	Me	NH2	2-Nap	Q-2			
Q-3	cPenO	H	NH2	2-Nap	Q-3	A	4.78	376(M <sup>+</sup> +1)
Q-4	cPenO	Me	NO2	5-Ind	Q-4			
Q-5	cPenO	Me	NH2	5-Ind	Q-5			
Q-6	cPenO	H	NH2	5-Ind	Q-6	A	3.75	365(M <sup>+</sup> +1)
Q-7	cPenO	H	NH2	1Me-5-Ind	Q-4,Q-5,Q-6	A	4.19	379(M <sup>+</sup> +1)
Q-8	cPenO	Me	NO2	1Me-5-1HIdz	Q-8			
Q-9	cPenO	Me	NH2	1Me-5-1HIdz	Q-9			
Q-10	cPenO	H	NH2	1Me-5-1HIdz	Q-10			
Q-11	cPenO	H	NH2	5-1HIdz	Q-8,Q-9,Q-10			
Q-12	cPenO	H	NH2	5-Bzt	Q-8,Q-9,Q-10			
Q-13	cPenO	H	NH2	5-2ABzt	Q-8,Q-9,Q-10			
Q-14	cPenO	H	NH2	2Me-5-Bzt	Q-8,Q-9,Q-10			
Q-15	cHexO	H	NH2	2-Nap	Q-1,Q-2,Q-3	A	5.66	404(M <sup>+</sup> +1)
Q-16	cHexO	H	NH2	1Me-5-Ind	Q-4,Q-5,Q-6			
Q-17	cHexO	H	NH2	1Me-5-1HIdz	Q-8,Q-9,Q-10			
Q-18	2EtBuO	H	NH2	2-Nap	Q-1,Q-2,Q-3			
Q-19	2EtBuO	H	NH2	5-Ind	Q-4,Q-5,Q-6	A	4.26	381(M <sup>+</sup> +1)
Q-20	2EtBuO	H	NH2	1Me-5-Ind	Q-4,Q-5,Q-6			
Q-21	2EtBuO	H	NH2	5-1HIdz	Q-8,Q-9,Q-10			
Q-22	2EtBuO	H	NH2	1Me-5-1HIdz	Q-8,Q-9,Q-10			
Q-23	2EtBuO	H	NH2	5-Bzt	Q-8,Q-9,Q-10			
Q-24	2EtBuO	H	NH2	5-2ABzt	Q-8,Q-9,Q-10			
Q-25	2EtBuO	H	NH2	2Me-5-Bzt	Q-8,Q-9,Q-10			
Q-26	iBuO	H	NH2	2-Nap	Q-1,Q-2,Q-3	A	4.82	364(M <sup>+</sup> +1)
Q-27	iBuO	H	NH2	1Me-5-Ind	Q-4,Q-5,Q-6			
Q-28	iBuO	H	NH2	1Me-5-1HIdz	Q-8,Q-9,Q-10	A	3.66	368(M <sup>+</sup> +1)
Q-29	(S)1PhEtO	H	NH2	2-Nap	Q-1,Q-2,Q-3	A	4.87	412(M <sup>+</sup> +1)
Q-30	(S)1PhEtO	H	NH2	1Me-5-Ind	Q-4,Q-5,Q-6	A	4.31	415(M <sup>+</sup> +1)
Q-31	(S)1PhEtO	H	NH2	1Me-5-1HIdz	Q-8,Q-9,Q-10	A	3.76	416(M <sup>+</sup> +1)
Q-32	4CF <sub>3</sub> BnO	H	NH2	2-Nap	Q-1,Q-2,Q-3	A	5.26	466(M <sup>+</sup> +1)
Q-33	4CF <sub>3</sub> BnO	H	NH2	1Me-5-Ind	Q-4,Q-5,Q-6	A	4.20	455(M <sup>+</sup> +1)
Q-34	4CF <sub>3</sub> BnO	H	NH2	1Me-5-1HIdz	Q-8,Q-9,Q-10			
Q-35	2-IndanO	H	NH2	2-Nap	Q-1,Q-2,Q-3	A	5.10	424(M <sup>+</sup> +1)
Q-36	2-IndanO	H	NH2	1Me-5-Ind	Q-4,Q-5,Q-6	A	4.63	427(M <sup>+</sup> +1)
Q-37	2-IndanO	H	NH2	1Me-5-1HIdz	Q-8,Q-9,Q-10	A	4.14	428 (M <sup>+</sup> +1)
Q-38	5OMe-2-IndanO	H	NH2	2-Nap	Q-1,Q-2,Q-3			
Q-39	5,8(OMe)-2-IndanO	H	NH2	1Me-5-Ind	Q-4,Q-5,Q-6			
Q-40	5F-2-IndanO	H	NH2	1Me-5-1HIdz	Q-8,Q-9,Q-10			
Q-41	2(4FPh)EtO	H	NH2	2-Nap	Q-1,Q-2,Q-3			
Q-42	2(4FPh)EtO	H	NH2	1Me-5-Ind	Q-4,Q-5,Q-6			
Q-43	2(4FPh)EtO	H	NH2	1Me-5-1HIdz	Q-8,Q-9,Q-10	A	4.48	448(M <sup>+</sup> +1)
Q-44	2(4DMAPh)EtO	H	NH2	2-Nap	Q-1,Q-2,Q-3	A	4.28	455(M <sup>+</sup> +1)
Q-45	2(4DMAPh)EtO	H	NH2	1Me-5-Ind	Q-4,Q-5,Q-6			
Q-46	2(4DMAPh)EtO	H	NH2	1Me-5-1HIdz	Q-8,Q-9,Q-10	A	3.12	459(M <sup>+</sup> +1)
Q-47	BnO	Me	NO2	2-Nap	Q-47			
Q-48	BnO	Me	NH2	2-Nap	Q-48			
Q-49	BnO	H	NH2	2-Nap	Q-3			
Q-50	BnO	Me	NO2	1Me-5-1HIdz	Q-47			
Q-51	BnO	Me	NH2	1Me-5-1HIdz	Q-48			
Q-52	BnO	H	NH2	1Me-5-1HIdz	Q-10			

[Example S-1]

Synthesis of methyl 3-{4-benzyloxy-3-(naphthalen-2-yl)-5-[N-(2,2,2-trifluoroacetyl)aminolphenyl}propionate (Intermediate 52)

According to the procedure described in the synthesis method of Compound No. B-103 with the modifications that the reaction was carried out for 1.5 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 4:1), Compound No. Q-48 (4.18 g), triethylamine (4.65 ml) and trifluoroacetic anhydride (7.40 ml) were reacted and treated to obtain the title compound (Intermediate 52, 4.72 g).

Synthesis of methyl 3-{4-hydroxy-3-(naphthalen-2-yl)-5-[N-(2,2,2-trifluoroacetyl)aminolphenyl}propionate (Intermediate 53)

A solution of Intermediate 52 (3.20 g) in a mixture of ethyl acetate (50 ml) and methanol (25 ml) was added with 10% palladium/carbon (98 mg), and stirred at room temperature for 2 hours under hydrogen atmosphere. The reaction mixture was filtered, and the solvent of the filtrate was evaporated under reduced pressure to obtain the title compound (Intermediate 53, 2.39 g).

Synthesis of methyl 3-{4-cyclopentyloxy-3-(naphthalen-2-yl)-5-[N-(2,2,2-trifluoroacetyl)aminolphenyl}propionate (Intermediate 54)

According to the procedure described in the synthesis method of Compound No. A-6 with the modifications that the reaction was carried out for 15.5 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 19:1), Intermediate 53 (84 mg), Ph<sub>3</sub>P (125 mg), cyclopentanol (50  $\mu$ l) and 40% DIAD (224  $\mu$ l) were reacted and treated to obtain the title compound (Intermediate 54, 90 mg).

Synthesis of methyl 3-{4-cyclopentyloxy-3-[N-methyl-N-(2,2,2-trifluoroacetyl)amino]-5-(naphthalen-2-yl)phenyl}propionate (Intermediate 55)

A solution of Intermediate 54 (208 mg) in DMF (5 ml) was added with 60%

sodium hydride (21 mg) under ice cooling, and stirred for 20 minutes. This reaction mixture was added dropwise with methyl iodide (150  $\mu$ l), stirred for 10 minutes, then warmed to room temperature, and further stirred for 1 hour. The reaction mixture was poured into ice water, and ethyl acetate (100 ml) was added for extraction. The organic layer was successively washed with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride, and saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 5:1) to obtain the title compound (Intermediate 55, 200 mg).

Synthesis of 3-[4-cyclopentyloxy-3-(N-methylamino)-5-(naphthalen-2-yl)phenyl]propionic acid (Compound No. S-1)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 6 hours, Intermediate 55 (198 mg) and 2 N aqueous sodium hydroxide (800  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. S-1, 38 mg).

[Example S-3]

Synthesis of methyl 3-[3-acetylamino-4-cyclopentyloxy-5-(naphthalen-2-yl)phenyl]propionate (Compound No. S-3)

A solution of Compound No. Q-2 (81 mg) in methylene chloride (2 ml) was added with N-methylmorpholine (33  $\mu$ l, WAKO), and added with acetyl chloride (22  $\mu$ l) under ice cooling. The reaction mixture was stirred for 10 minutes, then warmed to room temperature, and further stirred for 18 hours. The reaction mixture was poured into aqueous sodium hydrogencarbonate (100 ml), and added with ethyl acetate (150 ml) for extraction. The organic layer was successively washed with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride, and saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column

chromatography (Quad, hexane:ethyl acetate = 6:1) to obtain the title compound (Compound No. S-3, 85 mg).

[Example S-4]

Synthesis of 3-[3-acetylamino-4-cyclopentylmethoxy-5-(naphthalen-2-yl)phenyl]propionic acid (Compound No. S-4)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 15 hours, Compound No. S-3 (80 mg) and 2 N aqueous sodium hydroxide (400  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. S-4, 75 mg).

[Example S-5]

Synthesis of 3-[4-cyclopentylloxy-3-formylamino-5-(naphthalen-2-yl)phenyl]propionic acid (Compound No. S-5)

A solution of Compound No. Q-2 (90 mg) in DMF (5 ml) was added with a mixture of formic acid (200  $\mu$ l) and acetic anhydride (100  $\mu$ l) under ice cooling. The reaction mixture was stirred 10 minutes, then warmed to room temperature, and further stirred for 18 hours. The reaction mixture was poured into aqueous sodium hydrogencarbonate (100 ml), and added with ethyl acetate (150 ml) for extraction. The organic layer was successively washed with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride, and saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 5:1). The obtained substance was reacted and treated with 2N aqueous sodium hydroxide (400  $\mu$ l) according to the procedure described in the synthesis method of Intermediate 9 to obtain the title compound (Compound No. S-5, 65 mg).

[Example S-6]

Synthesis of methyl 3-[3-(2-acetoxyacetylamino)-4-cyclopentylloxy-5-(naphthalen-2-yl)phenyl]propionate (Compound No. S-6)



According to the procedure described in the synthesis method of Intermediate 70, Compound No. Q-2 (88 mg), N-methylmorpholine (36  $\mu$ l) and acetoxyacetyl chloride (35  $\mu$ l, Ald) were reacted and treated to obtain the title compound (Compound No. S-6, 75 mg).

[Example S-7]

Synthesis of 3-[4-cyclopentyloxy-3-(2-hydroxyacetyl-amino)-5-(naphthalen-2-yl)phenyl]propionic acid (Compound No. S-7)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 15.5 hours, Compound No. S-6 (102 mg) and 2 N aqueous sodium hydroxide (500  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. S-7, 80 mg).

[Example S-8]

Synthesis of 3-[3-carbamoylamino-4-cyclopentyloxy-5-(naphthalen-2-yl)phenyl]propionic acid (Compound No. S-8)

A solution of Compound No. Q-2 (100 mg) in a mixture of acetic acid (2 ml) and purified water (0.4 ml) was added with potassium cyanate (45 mg, Wako Pure Chemical Industries), and stirred at room temperature for 1 hour. The reaction mixture was poured into water (50 ml) containing ice, and extracted with isopropyl ether (150 ml x 2). The organic layer was successively washed with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride, and saturated brine and dried, and then the solvent was evaporated under reduced pressure. The obtained substance was reacted with 2 N aqueous sodium hydroxide (300  $\mu$ l) and treated according to the procedure described in the synthesis method of Intermediate 9 to obtain the title compound (Compound No. S-8, 70 mg).

[Example S-9]

Synthesis of methyl 3-[4-cyclopentyloxy-3-methylsulfonylamino-5-(naphthalen-2-yl)phenyl]propionate (Compound No. S-9)

A solution of Compound No. Q-2 (81 mg) in methylene chloride (2 ml) was added with pyridine (300  $\mu$ l), and then added with methanesulfonyl chloride (40  $\mu$ l) under ice cooling. The reaction mixture was stirred for 10 minutes, then warmed to room temperature, and further stirred for 2 hours. The reaction mixture was poured into 1 N hydrochloric acid, and added with ethyl acetate (150 ml) for extraction. The organic layer was washed successively with saturated aqueous sodium hydrogencarbonate, and saturated brine, and dried, and then the solvent of the organic layer was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 13:2) to obtain the title compound (Compound No. S-9, 96 mg).

Synthesis of 3-[4-cyclopentyloxy-3-methylsulfonylamino-5-(naphthalen-2-yl)phenyl]propionic acid (Compound No. S-10)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out at room temperature for 17.5 hours and at 60°C for 3 hours, Compound No. S-9 (81 mg) and 2 N aqueous sodium hydroxide (400  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. S-10, 80 mg).

[Example S-11]

Synthesis of 3-[4-cyclopentyloxy-3-(N,N-dimethylsulfamoylamino)-5-(naphthalen-2-yl)phenyl]propionic acid (Compound No. S-11)

A solution of Compound No. Q-2 (163 mg) in pyridine (5 ml) was successively added with 4-dimethylaminopyridine (104 mg, TCI) and dimethylsulfamoyl chloride (520  $\mu$ l, TCI), and stirred for 5 days. The reaction mixture was added with water (30 ml) and ethyl acetate (90 ml) for extraction. The organic layer was washed with saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 6:1). The obtained substance was

reacted with 2 N aqueous sodium hydroxide (300  $\mu$ l) and treated according to the procedure described in the synthesis method of Intermediate 9 to obtain the title compound (Compound No. S-11, 105 mg).

[Example S-12]

Synthesis of 3-[4-cyclopentyloxy-3-(N,N-dimethylamino)-5-(naphthalen-2-yl)phenyl]propionic acid (Compound No. S-12)

A solution of Compound No. Q-2 (60 mg) in DMF (3 ml) was added with 60% sodium hydride (26 mg) under ice cooling, and stirred for 10 minutes. The reaction mixture was added with methyl iodide (100  $\mu$ l), stirred for 10 minutes, then warmed to 60°C, and further stirred for 2 hours. The reaction mixture was poured into water (20 ml), and ethyl acetate (50 ml) was added for extraction. The organic layer was successively washed with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride, and saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 8:1). The obtained substance was reacted with 2 N aqueous sodium hydroxide (150  $\mu$ l) and treated according to the procedure described in the synthesis method of Intermediate 9 to obtain the title compound (Compound No. S-12, 46 mg).

Synthesis of methyl 3-{4-benzyloxy-3-(1-methyl-1H-indazol-5-yl)-5-[N-(2,2,2-trifluoroacetyl)aminolphenyl]propionate (Intermediate 56)

According to the procedure described in the synthesis method of Compound No. B-103 with the modifications that the reaction was carried out for 1.5 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 3:1), Compound No. Q-51 (2.09 g), triethylamine (3.70 ml) and trifluoroacetic anhydride (2.35 ml) were reacted and treated to obtain the title compound (Intermediate 56, 2.36 g).

Synthesis of methyl 3-{4-hydroxy-3-(1-methyl-1H-indazol-5-yl)-5-[N-(2,2,2-

trifluoroacetyl)aminophenyl}propionate (Intermediate 57)

A solution of Intermediate 56 (1.62 g) in a mixture of ethyl acetate (10 ml) and methanol (3 ml) was added with 10% palladium/carbon (29 mg), and stirred at room temperature for 17 hours under hydrogen atmosphere. The reaction mixture was filtered, and the solvent of the filtrate was evaporated under reduced pressure to obtain the title compound (Intermediate 57, 1.19 g).

[Examples S-1 to S-73]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification including the examples described above are shown in Table-S-1 and Table-S-2.

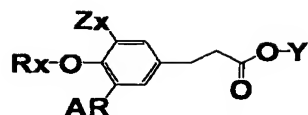


Table-S-1

Exp.	RxO	Y	Zx	AR	Syn	LCMS		
						method	RTime	Mass
S-1	cPenO	H	NHMe	2-Nap	S-1			
S-2	cPenO	H	NHEt	2-Nap	S-1			
S-3	cPenO	Me	NHAc	2-Nap	S-3			
S-4	cPenO	H	NHAc	2-Nap	S-4	C		421(M <sup>+</sup> +1)
S-5	cPenO	H	NHCHO	2-Nap	S-5	C		407(M <sup>+</sup> +1)
S-6	cPenO	H	NHCOCH <sub>2</sub> OAc	2-Nap	S-6			
S-7	cPenO	H	NHCOCH <sub>2</sub> OH	2-Nap	S-7	C		437(M <sup>+</sup> +1)
S-8	cPenO	H	NHCONH <sub>2</sub>	2-Nap	S-8	C		422(M <sup>+</sup> +1)
S-9	cPenO	Me	NHSO <sub>2</sub> Me	2-Nap	S-9			
S-10	cPenO	H	NHSO <sub>2</sub> Me	2-Nap	S-10	C		456(M <sup>+</sup> )
S-11	cPenO	H	NHSO <sub>2</sub> NMe <sub>2</sub>	2-Nap	S-11	C		483(M <sup>+</sup> +1)
S-12	cPenO	H	NMe <sub>2</sub>	2-Nap	S-12			
S-13	cPenO	H	NHMe	1Me-5-Ind	S-1			
S-14	cPenO	H	NMe <sub>2</sub>	1Me-5-Ind	S-12	C		407(M <sup>+</sup> +1)
S-15	cPenO	H	NHMe	1Me-5-1HIdz	S-1	C		394(M <sup>+</sup> +1)
S-16	cPenO	H	NMe <sub>2</sub>	1Me-5-1HIdz	S-12			
S-17	cPenO	H	NHMe	5-Bzt	S-1			
S-18	cPenO	H	NMe <sub>2</sub>	5-Bzt	S-12			
S-19	cPenO	H	NHMe	5-2ABzt	S-1			
S-20	cPenO	H	NMe <sub>2</sub>	5-2ABzt	S-12			
S-21	cPenO	H	NHMe	2Me-5-Bzt	S-1			
S-22	cPenO	H	NMe <sub>2</sub>	2Me-5-Bzt	S-12			
S-23	cPenMeO	H	NHMe	1Me-5-Ind	S-1			
S-24	cPenMeO	H	NMe <sub>2</sub>	1Me-5-Ind	S-12			
S-25	cPenMeO	H	NHMe	1Me-5-1HIdz	S-1			
S-26	cPenMeO	H	NMe <sub>2</sub>	1Me-5-1HIdz	S-12			
S-27	cHexO	H	NHMe	2-Nap	S-1			
S-28	cHexO	H	NMe <sub>2</sub>	2-Nap	S-12			
S-29	cHexO	H	NHMe	1Me-5-Ind	S-1	C		421(M <sup>+</sup> +1)
S-30	cHexO	H	NMe <sub>2</sub>	1Me-5-Ind	S-12			
S-31	cHexO	H	NHMe	1Me-5-1HIdz	S-1			
S-32	cHexO	H	NMe <sub>2</sub>	1Me-5-1HIdz	S-12			
S-33	2EtBuO	H	NHMe	2-Nap	S-1	C		406(M <sup>+</sup> +1)
S-34	2EtBuO	H	NHMe	6-OMe-2-Nap	S-1			
S-35	2EtBuO	H	NHMe	1Me-5-Ind	S-1			
S-36	2EtBuO	H	NHMe	5-Bzt	S-1			
S-37	2EtBuO	H	NHMe	1Me-5-1HIdz	S-1			
S-38	iBuO	H	NHMe	2-Nap	S-1			
S-39	iBuO	H	NMe <sub>2</sub>	2-Nap	S-12	C		392(M <sup>+</sup> +1)
S-40	iBuO	H	NHMe	1Me-5-Ind	S-1	C		381(M <sup>+</sup> +1)
S-41	iBuO	H	NMe <sub>2</sub>	1Me-5-Ind	S-12			
S-42	iBuO	H	NHMe	1Me-5-1HIdz	S-1			
S-43	iBuO	H	NMe <sub>2</sub>	1Me-5-1HIdz	S-12			
S-44	1PhEtO	H	NHMe	2-Nap	S-1	C		426(M <sup>+</sup> +1)
S-45	1PhEtO	H	NMe <sub>2</sub>	2-Nap	S-12			

Table-S-2

S-46	1PhEtO	H	NHMe	1Me-5-Ind	S-1			
S-47	1PhEtO	H	NMe <sub>2</sub>	1Me-5-Ind	S-12	C		443(M <sup>+</sup> +1)
S-48	1PhEtO	H	NHMe	1Me-5-1HIdz	S-1	C		429(M <sup>+</sup> +1)
S-49	1PhEtO	H	NMe <sub>2</sub>	1Me-5-1HIdz	S-12			
S-50	4CF <sub>3</sub> BnO	H	NHMe	2-Nap	S-1			
S-51	4CF <sub>3</sub> BnO	H	NMe <sub>2</sub>	2-Nap	S-12			
S-52	4CF <sub>3</sub> BnO	H	NHMe	1Me-5-Ind	S-1			
S-53	4CF <sub>3</sub> BnO	H	NMe <sub>2</sub>	1Me-5-Ind	S-12	C		497(M <sup>+</sup> +1)
S-54	4CF <sub>3</sub> BnO	H	NHMe	1Me-5-1HIdz	S-1			
S-55	4CF <sub>3</sub> BnO	H	NMe <sub>2</sub>	1Me-5-1HIdz	S-12			
S-56	2-IndanO	H	NHMe	2-Nap	S-1			
S-57	2-IndanO	H	NMe <sub>2</sub>	2-Nap	S-12			
S-58	2-IndanO	H	NHMe	1Me-5-Ind	S-1	C		441(M <sup>+</sup> +1)
S-59	2-IndanO	H	NMe <sub>2</sub>	1Me-5-Ind	S-12			
S-60	2-IndanO	H	NHMe	1Me-5-1HIdz	S-1	A	4.16	442(M <sup>+</sup> +1)
S-61	2-IndanO	H	NMe <sub>2</sub>	1Me-5-1HIdz	S-12	A	4.18	456(M <sup>+</sup> +1)
S-62	2(4FPh)EtO	H	NHMe	2-Nap	S-1			
S-63	2(4FPh)EtO	H	NMe <sub>2</sub>	2-Nap	S-12	C		458(M <sup>+</sup> +1)
S-64	2(4FPh)EtO	H	NHMe	1Me-5-Ind	S-1	C		447(M <sup>+</sup> +1)
S-65	2(4FPh)EtO	H	NMe <sub>2</sub>	1Me-5-Ind	S-12			
S-66	2(4FPh)EtO	H	NHMe	1Me-5-1HIdz	S-1			
S-67	2(4FPh)EtO	H	NMe <sub>2</sub>	1Me-5-1HIdz	S-12			
S-68	2(4DMAPh)EtO	H	NHMe	2-Nap	S-1	C		469(M <sup>+</sup> +1)
S-69	2(4DMAPh)EtO	H	NMe <sub>2</sub>	2-Nap	S-12			
S-70	2(4DMAPh)EtO	H	NHMe	1Me-5-Ind	S-1			
S-71	2(4DMAPh)EtO	H	NMe <sub>2</sub>	1Me-5-Ind	S-12	C		486(M <sup>+</sup> +1)
S-72	2(4DMAPh)EtO	H	NHMe	1Me-5-1HIdz	S-1	C		473(M <sup>+</sup> +1)
S-73	2(4DMAPh)EtO	H	NMe <sub>2</sub>	1Me-5-1HIdz	S-12			

## [Example T-1]

Synthesis of 3-[4-cyclopentylmethoxy-3-hydroxy-5-(naphthalen-2-yl)phenyl]propionic acid (Compound No. T-1)

A solution of Compound No. Q-2 (403 mg) in acetic acid (1.5 ml) was added with 20% sulfuric acid (1.0 ml). This reaction mixture was added dropwise with an aqueous solution (0.5 ml) of sodium nitrite (76 mg) over 10 minutes while keeping the temperature of the reaction mixture below 10°C, and further stirred for 5 minutes. This reaction solution was added to a solution of sodium acetate (328 mg) in acetic acid (3.5 ml) heated and stirred at 100°C beforehand, and further stirred for 10 minutes with heating. The reaction solution was poured into ice

water (50 ml), and extracted with isopropyl ether (100 ml x 2). The organic layer was washed successively with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride and saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 10:1). The obtained substance was reacted with 2 N aqueous sodium hydroxide (500  $\mu$ l) and treated according to the procedure described in the synthesis method of Intermediate 9 to obtain the title compound (Compound No. T-1, 78 mg).

[Example T-2]

Synthesis of ethyl 3-[3-acetoxy-4-cyclopentyloxy-5-(naphthalen-2-yl)phenyl]propionate (Intermediate 58)

According to the procedure described in the synthesis method of Compound No. C-1 with the modifications that the reaction was carried out for 13 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 9:1), Compound No. B-114 (160 mg), 2-naphthaleneboronic acid (382 mg, Ald), 2 M aqueous sodium carbonate (0.7 ml) and (Ph<sub>3</sub>P)<sub>4</sub>Pd (105 mg) were reacted and treated to obtain the title compound (Intermediate 58, 152 mg).

Synthesis of 3-[4-cyclopentyloxy-3-hydroxy-5-(naphthalen-2-yl)phenyl]propionic acid (Compound No. T-2)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 2 hours, Intermediate 58 (146 mg) and 2 N aqueous sodium hydroxide (0.35 ml) were reacted and treated to obtain the title compound (Compound No. T-2, 135 mg).

[Example T-31]

Synthesis of ethyl 3-[4-cyclopentyloxy-3-methoxy-5-(naphthalen-2-yl)phenyl]propionate (Compound No. T-31)

According to the procedure described in the synthesis method of Compound

No. C-1 with the modifications that the reaction was carried out for 14 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 9:1), Compound No. A-25 (210 mg), 2-naphthaleneboronic acid (184 mg), 2 M aqueous sodium carbonate (0.5 ml) and  $(\text{Ph}_3\text{P})_4\text{Pd}$  (65.3 mg) were reacted and treated to obtain the title compound (Compound No. T-31, 181 mg).

[Example T-32]

Synthesis of 3-[4-cyclopentyloxy-3-methoxy-5-(naphthalen-2-yl)phenyl]propionic acid (Compound No. T-32)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 2 hours, Compound No. T-31 (166 mg) and 2 N aqueous sodium hydroxide (0.45 ml) were reacted and treated to obtain the title compound (Compound No. T-32, 135 mg).

[Example T-33]

Synthesis of 4-(t-butyldimethylsilyloxy)-3-(1H-indol-5-yl)-5-methoxybenzaldehyde (Intermediate 59)

According to the procedure described in the synthesis method of Compound No. C-1 with the modifications that the reaction was carried out for 12.5 hours, and the purification was performed by flash column chromatography (hexane:ethyl acetate = 7:1), 5-indoleboronic acid (1.29 g), Intermediate 16 (1.75 g), 2 M aqueous sodium carbonate (4.8 ml) and  $(\text{Ph}_3\text{P})_4\text{Pd}$  (400 mg) were reacted and treated to obtain the title compound (Intermediate 59, 910 mg).

Synthesis of ethyl 3-[4-(t-butyldimethylsilyloxy)-3-(1H-indol-5-yl)-5-methoxyphenyl]acrylate (Intermediate 60)

According to the procedure described in the synthesis method of Intermediate 7 with the modifications that the reaction was carried out for 1.5 hours, and the purification was performed by flash column chromatography (hexane:ethyl acetate = 3:1), Intermediate 59 (910 mg), ethyl



diethylphosphonoacetate (500  $\mu$ l) and 60% sodium hydride (100 mg) were reacted and treated to obtain the title compound (Intermediate 60, 945 mg).

Synthesis of ethyl 3-[4-(t-butyldimethylsilyloxy)-3-(1H-indol-5-yl)-5-methoxyphenyl]propionate (Intermediate 61)

According to the procedure described in the synthesis method of Intermediate 8, Intermediate 60 (945 mg) and 10% palladium/carbon (95 mg) were reacted and treated under hydrogen gas atmosphere to obtain the title compound (Intermediate 61, 940 mg).

Synthesis of ethyl 3-[4-hydroxy-3-(1H-indol-5-yl)-5-methoxyphenyl]propionate (Intermediate 62)

According to the procedure described in the synthesis method of Intermediate 19 with the modifications that the reaction was carried out for 1.5 hours, and the purification was performed by flash column chromatography (hexane:ethyl acetate = 2:1), Intermediate 61 (750 mg) and a 1 M solution of tetrabutylammonium fluoride in THF (5.0 ml) were reacted and treated to obtain the title compound (Intermediate 62, 555 mg).

Synthesis of ethyl 3-[4-cyclopentyloxy-3-(1H-indol-5-yl)-5-methoxyphenyl]propionate (Compound No. T-33)

According to the procedure described in the synthesis method of Compound No. A-6 with the modifications that the reaction was carried out for 16 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 7:1), Intermediate 62 (340 mg), Ph<sub>3</sub>P (1.31 g), cyclopentanol (450  $\mu$ l) and TMAD (860 mg) were reacted and treated to obtain the title compound (Compound No. T-33, 376 mg).

[Example T-34]

Synthesis of 3-[4-cyclopentyloxy-3-(1H-indol-5-yl)-5-methoxyphenyl]propionic acid (Compound No. T-34)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 2 hours, Compound No. T-33 (99 mg) and 2 N aqueous sodium hydroxide (500  $\mu$  l) were reacted and treated to obtain the title compound (Compound No. T-34, 76 mg).

[Examples T-1 to T-61]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification including the examples described above are shown in Table-T-1 and Table T-2.

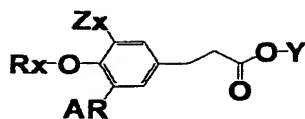


Table-T-1

Exp.	RxO	Y	Zx	AR	Syn	LCMS		
						method	RTime	Mass
T-1	cPenMeO	H	OH	2-Nap	T-1	A	5.03	382(M <sup>+</sup> +1)
T-2	cPenO	H	OH	2-Nap	T-2			
T-3	cPenO	H	OH	5-Ind	Int73,T-2	C		366(M <sup>+</sup> +1)
T-4	cPenO	H	OH	1Me-5-Ind	Int73,T-2			
T-5	cPenO	H	OH	5-1HIdz	Int73,T-2			
T-6	cPenO	H	OH	1Me-5-Idz	Int73,T-2	C		381(M <sup>+</sup> +1)
T-7	cHexO	H	OH	2-Nap	T-1			
T-8	cHexO	H	OH	1Me-5-Ind	T-1			
T-9	cHexO	H	OH	1Me-5-Idz	T-1			
T-10	2EtBuO	H	OH	2-Nap	T-1	C		393(M <sup>+</sup> +1)
T-11	2EtBuO	H	OH	1Me-5-Ind	T-1			
T-12	2EtBuO	H	OH	1Me-5-Idz	T-1			
T-13	iBuO	H	OH	2-Nap	T-1			
T-14	iBuO	H	OH	1Me-5-Ind	T-1			
T-15	iBuO	H	OH	1Me-5-Idz	T-1			
T-16	1PhEtO	H	OH	2-Nap	T-1			
T-17	1PhEtO	H	OH	1Me-5-Ind	T-1	C		416(M <sup>+</sup> +1)
T-18	1PhEtO	H	OH	1Me-5-Idz	T-1			
T-19	4CF <sub>3</sub> BnO	H	OH	2-Nap	T-1			
T-20	4CF <sub>3</sub> BnO	H	OH	1Me-5-Ind	T-1			
T-21	4CF <sub>3</sub> BnO	H	OH	1Me-5-Idz	T-1			
T-22	2-IndanO	H	OH	2-Nap	T-1			
T-23	2-IndanO	H	OH	1Me-5-Ind	T-1			
T-24	2-IndanO	H	OH	1Me-5-Idz	T-1	A	3.91	429(M <sup>+</sup> +1)
T-25	2(4FPh)EtO	H	OH	2-Nap	T-1			
T-26	2(4FPh)EtO	H	OH	1Me-5-Ind	T-1			
T-27	2(4FPh)EtO	H	OH	1Me-5-Idz	T-1			
T-28	2(4DMAPh)EtO	H	OH	2-Nap	T-1			
T-29	2(4DMAPh)EtO	H	OH	1Me-5-Ind	T-1	C		459(M <sup>+</sup> +1)
T-30	2(4DMAPh)EtO	H	OH	1Me-5-Idz	T-1			
T-31	cPenO	Et	OMe	2-Nap	T-31			
T-32	cPenO	H	OMe	2-Nap	T-32			
T-33	cPenO	Et	OMe	5-Ind	T-33			
T-34	cPenO	H	OMe	5-Ind	T-34			
T-35	cPenO	H	OMe	1Me-5-Ind	T-33,T-34	A	4.72	394(M <sup>+</sup> +1)
T-36	cPenO	H	OMe	5-1HIdz	T-31,T-32			
T-37	cPenO	H	OMe	1Me-5-Idz	T-31,T-32			
T-38	cHexO	H	OMe	2-Nap	T-31,T-32	C		405(M <sup>+</sup> +1)
T-39	cHexO	H	OMe	1Me-5-Ind	T-33,T-34			
T-40	cHexO	H	OMe	1Me-5-Idz	T-31,T-32			
T-41	2EtBuO	H	OMe	2-Nap	T-31,T-32			
T-42	2EtBuO	H	OMe	1Me-5-Ind	T-33,T-34			
T-43	2EtBuO	H	OMe	1Me-5-Idz	T-31,T-32			

Table-T-2

T-44	iBuO	H	OMe	2-Nap	T-31,T-32			
T-45	iBuO	H	OMe	1Me-5-Ind	T-33,T-34	C		382(M <sup>+</sup> +1)
T-46	iBuO	H	OMe	1Me-5-1HIdz	T-31,T-32			
T-47	1PhEtO	H	OMe	2-Nap	T-31,T-32			
T-48	1PhEtO	H	OMe	1Me-5-Ind	T-33,T-34			
T-49	1PhEtO	H	OMe	1Me-5-1HIdz	T-31,T-32	C		431(M <sup>+</sup> +1)
T-50	4CF <sub>3</sub> BnO	H	OMe	2-Nap	T-31,T-32			
T-51	4CF <sub>3</sub> BnO	H	OMe	1Me-5-Ind	T-33,T-34			
T-52	4CF <sub>3</sub> BnO	H	OMe	1Me-5-1HIdz	T-31,T-32			
T-53	2-IndanO	H	OMe	2-Nap	T-31,T-32			
T-54	2-IndanO	H	OMe	1Me-5-Ind	T-33,T-34			
T-55	2-IndanO	H	OMe	1Me-5-1HIdz	T-31,T-32	C		443(M <sup>+</sup> +1)
T-56	2(4FPh)EtO	H	OMe	2-Nap	T-31,T-32			
T-57	2(4FPh)EtO	H	OMe	1Me-5-Ind	T-33,T-34	C		448(M <sup>+</sup> +1)
T-58	2(4FPh)EtO	H	OMe	1Me-5-1HIdz	T-31,T-32			
T-59	2(4DMAPh)EtO	H	OMe	2-Nap	T-31,T-32	C		470(M <sup>+</sup> +1)
T-60	2(4DMAPh)EtO	H	OMe	1Me-5-Ind	T-33,T-34			
T-61	2(4DMAPh)EtO	H	OMe	1Me-5-1HIdz	T-31,T-32			

## [Example U-1]

Synthesis of 4-cyclohexylmethyloxy-3-(naphthalen-2-yl)phenylacetonitrile

(Intermediate 63)

A solution of Compound No. C-1 (172 mg) in dehydrated THF (5 ml) was added successively with trimethylsilylnitrile (133  $\mu$ l, TCI) under ice cooling and zinc iodide (16 mg, WAKO) under argon gas atmosphere, stirred for 15 minutes, then warmed to room temperature, and further stirred for 27 hours. The reaction mixture was added with ethyl acetate (90 ml), and washed successively with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride and saturated brine. The organic layer was dried, and then the solvent was evaporated under reduced pressure. A solution of the residue in anhydrous methylene chloride (5 ml) was added with triethylsilane (240  $\mu$ l, TCI) under ice cooling and boron trifluoride diethyl ether complex (366  $\mu$ l, TCI) under argon gas atmosphere, warmed to room temperature, and stirred for 3.5 hours. The reaction mixture was poured into ice water (50 ml), and extracted with ethyl acetate (90 ml). The organic layer was successively washed with saturated aqueous sodium

hydrogencarbonate, saturated aqueous ammonium chloride, and saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 10:1) to obtain the title compound (Intermediate 63, 116 mg).

Synthesis of 4-cyclohexylmethoxy-3-(naphthalen-2-yl)phenylacetic acid  
(Compound No. U-1)

According to the procedure described in the synthesis method of Intermediate 9 with the modifications that the reaction was carried out for 24 hours under reflux by heating, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 2:1), Intermediate 63 (110 mg) and 5 N aqueous sodium hydroxide (900  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. U-1, 62 mg).

[Example U-10]

Synthesis of methyl 4-[4-cyclopentylmethoxy-3-(naphthalen-2-yl)phenyl]butyrate  
(Compound No. U-10)

According to the procedure described in the synthesis method of Compound No. C-1 with the modifications that the reaction was carried out for 18 hours, and the purification was performed by column chromatography (Quad, hexane:isopropyl ether = 8:1), Compound No. F-1 (355 mg), 2-naphthaleneboronic acid (344 mg), 2 M aqueous sodium carbonate (2.1 ml) and  $(\text{Ph}_3\text{P})_4\text{Pd}$  (115 mg) were reacted and treated to obtain the title compound (Compound No. U-10, 392 mg).

[Example U-11]

Synthesis of 4-[4-cyclopentylmethoxy-3-(naphthalen-2-yl)phenyl]butyric acid  
(Compound No. U-11)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 3.5 hours, Compound No. U-10 (380 mg) and 2 N aqueous sodium hydroxide (1.0 ml) were reacted and

treated to obtain the title compound (Compound No. U-11, 342 mg).

[Examples U-1 to U-18]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification including the examples described above are shown in Table-U-1.

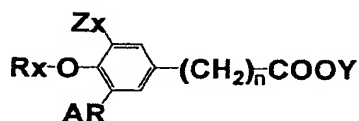


Table-U-1

Exp.	RxO	Y	Zx	n	AR	Syn	LCMS		
							method	RTime	Mass
U-1	cHexMeO	H	H	1	2-Nap	U-1	C		374(M <sup>+</sup> )
U-2	cHexMeO	H	H	1	1Me-5-Ind	Int63,U-1			
U-3	cHexMeO	H	H	1	1Me-5-Idz	Int63,U-1			
U-4	cPenMeO	H	H	1	2-Nap	Int63,U-1	C		360(M <sup>+</sup> )
U-5	cPenMeO	H	H	1	1Me-5-Ind	Int63,U-1			
U-6	cPenO	H	H	1	2-Nap	Int63,U-1			
U-7	cPenO	H	H	1	1Me-5-Ind	Int63,U-1	C		349(M <sup>+</sup> )
U-8	2(4FPh)EtO	H	H	1	2-Nap	Int63,U-1			
U-9	2(4FPh)EtO	H	H	1	1Me-5-Ind	Int63,U-1			
U-10	cPenMeO	Me	H	3	2-Nap	U-10	C		374(M <sup>+</sup> )
U-11	cPenMeO	H	H	3	2-Nap	U-11	C		374(M <sup>+</sup> )
U-12	cPenMeO	H	H	3	1Me-5-Ind	U-10,U-11			
U-13	cPenO	H	H	3	2-Nap	U-10,U-11			
U-14	cPenO	H	H	3	1Me-5-Ind	U-10,U-11	C		377(M <sup>+</sup> )
U-15	cHexO	H	H	3	2-Nap	U-10,U-11			
U-16	cHexO	H	H	3	1Me-5-Ind	U-10,U-11			
U-17	2(4FPh)EtO	H	H	3	2-Nap	U-10,U-11			
U-18	2(4FPh)EtO	H	H	3	1Me-5-Ind	U-10,U-11			

[Example V-1]

Synthesis of ethyl 3-[4-cyclohexylmethyloxy-3-(naphthalen-1-yl)phenyl]acrylate  
(Intermediate 64)

According to the procedure described in the synthesis method of Intermediate 7 provided that the reaction was carried out for 1 hour, Compound No. C-2 (361 mg), ethyl diethylphosphonoacetate (240  $\mu$ l), 60% sodium hydride (69 mg) were reacted and treated to obtain the title compound (Intermediate 64, 377 mg).  
Synthesis of ethyl 3-[4-cyclohexylmethyloxy-3-(naphthalen-1-yl)phenyl]propionate

(Compound No. V-1)

According to the procedure described in the synthesis method of Intermediate 8 with the modifications that the reaction was carried out for 1.5 hours, and the purification was performed by flash column chromatography (hexane:ethyl acetate = 10:1), Intermediate 64 (361 mg) and 10% palladium/carbon (49 mg) were reacted under hydrogen atmosphere and treated to obtain the title compound (Compound No. V-1, 344 mg).

[Example V-2]

Synthesis of 3-[4-cyclohexylmethyloxy-3-(naphthalen-1-yl)phenyl]propionic acid (Compound No. V-2)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 1.5 hours, Compound No. V-1 (332 mg) and 2 N aqueous sodium hydroxide (900  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. V-2, 295 mg).

[Example V-3]

Synthesis of methyl 3-[4-cyclopentylmethyloxy-3-(6-hydroxynaphthalen-2-yl)phenyl]propionate (Compound No. V-3)

A solution of 2-bromo-6-hydroxynaphthalene (243 mg, TCI) in anhydrous THF (10 ml) was cooled to  $-78^{\circ}\text{C}$ , added dropwise with a 1.6 M solution of n-butyllithium in hexane (1.18 ml) over 20 minutes under argon gas atmosphere, and stirred for 30 minutes. The reaction mixture was added dropwise with  $(\text{iPrO})_3\text{B}$  (1.73 ml) over 10 minutes, stirred for 30 minutes, then warmed to room temperature, and further stirred for 2 hours. The reaction mixture was added with 0.5 M aqueous sulfuric acid (2 ml), and extracted with diethyl ether (40 ml x 3). The organic layer was washed with saturated brine and dried, and then the solvent was evaporated under reduced pressure to obtain crude 6-hydroxy-2-naphthaleneboronic acid (378 mg). A solution of this substance in ethanol (1 ml),

Compound No. A-1 (230 mg), and 2 M aqueous sodium carbonate (2.4 ml) were added with toluene (3 ml) and  $(\text{Ph}_3\text{P})_4\text{Pd}$  (115 mg) and stirred at 100°C for 13 hours. The reaction mixture was added with ethyl acetate (100 ml), and washed successively with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride and saturated brine. The organic layer was dried, and then the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography (hexane:ethyl acetate = 6:1) to obtain the title compound (Compound No. V-3, 270 mg).

[Example V-4]

Synthesis of 3-[4-cyclopentylmethyloxy-3-(6-hydroxynaphthalen-2-yl)phenyl]propionic acid (Compound No. V-4)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 14 hours, Compound No. V-3 (149 mg) and 2 N aqueous sodium hydroxide (370  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. V-4, 117 mg).

[Example V-5]

Synthesis of methyl 3-[4-cyclopentylmethyloxy-3-(5-hydroxynaphthalen-2-yl)phenyl]propionate (Compound No. V-5)

2-Amino-5-hydroxynaphthalene (4.80 g, TCI) was dissolved in 6 N hydrochloric acid (300 ml), added dropwise with an aqueous solution (22.5 ml) of sodium nitrite (2.25 g) over 30 minutes under ice cooling, and stirred for 30 minutes. The reaction mixture was added dropwise with an aqueous solution (75 ml) of potassium iodide (9.90 g, WAKO), stirred for 30 minutes, then warmed to room temperature, and further stirred for 3.5 hours. The reaction mixture was neutralized with aqueous ammonia, and then filtered through a Celite layer. The filtrate was added with ethyl acetate (90 ml x 2) for extraction. The organic layer was washed successively with saturated aqueous sodium hydrogencarbonate,



saturated aqueous ammonium chloride and saturated brine, and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 10:1) to obtain 1-hydroxy-6-iodonaphthalene (1.48 g). A solution of this substance (539 mg) in anhydrous THF (10 ml) was added with 60% sodium hydride (171 mg) under ice cooling, and stirred for 1 hour. The reaction mixture was cooled to -78°C under argon gas atmosphere, added dropwise with a 1.6 M solution of n-butyllithium in hexane (3.75 ml) over 10 minutes, and stirred for 30 minutes. The reaction mixture was added dropwise with (iPrO)<sub>3</sub>B (1.16 ml) over 10 minutes, stirred for 30 minutes, then warmed to room temperature, and further stirred for 3 hours. The reaction mixture was added with water (3 ml) and 0.5 M aqueous sulfuric acid (7 ml), and extracted with diethyl ether (100 ml x 3). The organic layer was washed with saturated brine and dried, and then the solvent was evaporated under reduced pressure to obtain crude 7-hydroxy-2-naphthaleneboronic acid. A solution of this substance in ethanol (1 ml), Compound No. A-1 (350 mg), 2 M aqueous sodium carbonate (2.4 ml) and (Ph<sub>3</sub>P)<sub>4</sub>Pd (116 mg) were reacted and treated according to the procedure described in the synthesis method of Compound No. V-3 with the modifications that the reaction was carried out for 14 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 6:1) to obtain the title compound (Compound No. V-5, 388 mg).

#### [Example V-6]

Synthesis of 3-[4-cyclopentylmethoxy-3-(5-hydroxynaphthalen-2-yl)phenyl]propionic acid (Compound No. V-6)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 12 hours, Compound No. V-5 (355 mg) and 2 N aqueous sodium hydroxide (1.75 ml) were reacted and treated to obtain the title compound (Compound No. V-6, 158 mg).

[Example V-7]

Synthesis of methyl 3-[4-cyclopentylmethoxy-3-(7-hydroxynaphthalen-2-yl)phenyl]propionate (Compound No. V-7)

According to the procedure described in the synthesis method of Compound No. V-5 with the modifications that the reaction was carried out for 4 hours, and the purification was performed by flash column chromatography (hexane:ethyl acetate = 6:1), crude 7-hydroxy-2-naphthaleneboronic acid prepared from 2-bromo-7-hydroxynaphthalene (559 mg, MAYB), a 1.6M solution of n-butyllithium in hexane (3.91 ml) and (iPrO)<sub>3</sub>B (1.16 ml), Compound No. A-1 (386 mg), 2 M aqueous sodium carbonate (4.0 ml) and (Ph<sub>3</sub>P)<sub>4</sub>Pd (195 mg) were reacted and treated to obtain the title compound (Compound No. V-7, 460 mg).

[Example V-8]

Synthesis of 3-[4-cyclopentylmethoxy-3-(7-hydroxynaphthalen-2-yl)phenyl]propionic acid (Compound No. V-8)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 27 hours, Compound No. V-7 (176 mg) and 2 N aqueous sodium hydroxide (436  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. V-8, 109 mg).

[Example V-11]

Synthesis of methyl 3-{4-cyclohexylmethoxy-3-[6-(N,N-dimethylcarbamoylmethoxy)naphthalen-2-yl]phenyl}propionate (Compound No. V-11)

A solution of Compound No. V-3 (185 mg) in DMF (5 ml) was added with potassium carbonate (274 mg), and 2-chloro-N,N-dimethylacetamide (411  $\mu$ l, KANTO), and stirred at 50°C for 18 hours. The reaction mixture was added with ethyl acetate (90 ml), and washed with saturated brine. The organic layer was dried, and then the solvent was evaporated under reduced pressure. The residue

was purified by PTLC (chloroform:methanol = 10:1) to obtain the title compound (Compound No. V-11, 213 mg).

[Example V-12]

Synthesis of 3-{4-cyclohexylmethoxy-3-[6-(N,N-dimethylcarbamoylmethoxy)naphthalen-2-yl]phenyl}propionic acid (Compound No. V-10)

According to the procedure described in the synthesis method of Intermediate 9 with the modifications that the reaction was carried out at room temperature for 18 hours and at 60°C for 8 hours, and the purification was performed by PTLC (chloroform:methanol = 10:1), Compound No. V-11 (213 mg) and 2 N aqueous sodium hydroxide (420  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. V-12, 115 mg).

[Example V-13]

Synthesis of methyl 3-[3-(6-aminonaphthalen-2-yl)-4-cyclopentylmethoxyphenyl]propionate (Compound No. V-13)

According to a known method described in a publication (Anderson, L.C. et al., J. Am. Chem. Soc, 1943, vol. 65, p.241), a solution of 2-amino-6-bromonaphthalene (223 mg) obtainable from commercially available 2-bromo-6-hydroxynaphthalene (TCI) in anhydrous THF (10 ml) was added with 30% potassium hydride (191 mg, Ald) under ice cooling, and stirred for 1 hour. The reaction mixture was cooled to -78°C under argon gas atmosphere, added dropwise with a 1.7 M solution of t-butyllithium in pentane (1.88 ml) over 10 minutes, and stirred for 30 minutes. This reaction mixture was added dropwise with (iPrO)<sub>3</sub>B (0.92 ml) over 10 minutes, stirred for 30 minutes, then warmed to room temperature, and further stirred for 3 hours. The reaction mixture was added with water (3 ml) and 0.5 M aqueous sulfuric acid (4 ml), and extracted with diethyl ether (100 ml x 3). The organic layer was washed with saturated brine and dried,

and then the solvent was evaporated under reduced pressure to obtain crude 6-amino-2-naphthaleneboronic acid (402 mg). A solution of this substance in ethanol (0.5 ml), Compound No. A-1 (119 mg), 2 M aqueous sodium carbonate (1.5 ml) and  $(\text{Ph}_3\text{P})_4\text{Pd}$  (61 mg) were reacted and treated according to the procedure described in the synthesis method of Compound No. V-3 with the modifications that the reaction was carried out for 13 hours, and the purification was performed by flash column chromatography (hexane:ethyl acetate = 4:1) to obtain the title compound (Compound No. V-13, 129 mg).

[Example V-14]

Synthesis of 3-[3-(6-aminonaphthalen-2-yl)-4-cyclopentylmethyloxyphenyl]propionic acid (Compound No. V-14)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 14 hours, Compound No. V-13 (120 mg) and 2 N aqueous sodium hydroxide (1.75 ml) were reacted and treated to obtain the title compound (Compound No. V-14, 89 mg).

[Example V-16]

Synthesis of methyl 3-[3-({6-[2-(acetyloxy)acetylamino)naphthalen-2-yl]}-4-cyclopentylmethyloxyphenyl)propionate (Intermediate 65)

A solution of Compound No. V-13 (151 mg) in dichloromethane (4 ml) was added with N-methylmorpholine (50  $\mu$ l), and then added with acetyloxyacetyl chloride (48.3  $\mu$ l) under ice cooling. The reaction mixture was stirred for 10 minutes, then warmed to room temperature, and further stirred for 4 hours. The reaction mixture was poured into aqueous sodium hydrogencarbonate (100 ml), and ethyl acetate (150 ml) was added for extraction. The organic layer was successively washed with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride, and saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by PTLC

(hexane:ethyl acetate = 1:1) to obtain the title compound (Intermediate 88, 136 mg).  
Synthesis of 3-(4-cyclopentylmethoxy-3-(6-[2-(hydroxyacetyl)amino]naphthalen-2-yl)phenyl)propionic acid (Compound No. V-16)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out at room temperature for 5 hours and at 60°C for 1 hour, Intermediate 65 (135 mg) and 2 N aqueous sodium hydroxide (1.12 ml) were reacted and treated to obtain the title compound (Compound No. V-16, 102 mg).

[Example V-18]

Synthesis of methyl 3-[4-cyclopentylmethoxy-3-(6-methylsulfonylaminonaphthalen-2-yl)phenyl]propionate (Compound No. V-18)

A solution of Compound No. V-13 (149.1 mg) in 1,2-dichloroethane (5 ml) was added successively with pyridine (500  $\mu$ l) and methanesulfonyl chloride (62  $\mu$ l) under ice cooling, stirred for 1.5 hours, then warmed to room temperature, and stirred for 12 hours. The reaction mixture was added with water (30 ml) and ethyl acetate (90 ml) for extraction. The organic layer was successively washed with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride, and saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by PTLC (hexane:ethyl acetate = 2:1) to obtain the title compound (Compound No. V-18, 126 mg).

[Example V-19]

Synthesis of 3-[4-cyclopentylmethoxy-3-(6-methylsulfonylaminonaphthalen-2-yl)phenyl]propionic acid (Compound No. V-19)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out at room temperature for 3 hours and at 60°C for 1 hour, Compound No. V-18 (129 mg) and 2 N aqueous sodium hydroxide (535  $\mu$ l) were reacted and treated to obtain the title compound

(Compound No. V-19, 98 mg).

[Example V-20]

Synthesis of methyl 3-{4-cyclopentylmethoxy-3-[6-(N,N-dimethylsulfamoylamino)naphthalen-2-yl]phenyl}propionate (Compound No. V-20)

A solution of Compound No. V-13 (165 mg) in pyridine (5 ml) was added successively with 4-dimethylaminopyridine (104 mg, TCI) and dimethylsulfamoyl chloride (520  $\mu$ l, TCI), stirred for 5 days, and then further stirred at 50°C for 4 hours. The reaction mixture was added with water (30 ml) and ethyl acetate (90 ml)) for extraction. The organic layer was washed with saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 6:1) to obtain the title compound (Compound No. V-20, 125 mg).

[Example V-21]

Synthesis of 3-{4-cyclopentylmethoxy-3-[6-(N,N-dimethylsulfamoylamino)naphthalen-2-yl]phenyl}propionic acid (Compound No. V-21)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 1.5 hours, Compound No. V-20 (118 mg) and 2 N aqueous sodium hydroxide (460  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. V-21, 87 mg).

[Example V-22]

Synthesis of 2-bromo-6-sulfamoylaminonaphthalene (Intermediate 66)

A solution of chlorosulfonyl isocyanate (870  $\mu$ l, WAKO) in benzene (10 ml) was added dropwise with formic acid (377  $\mu$ l, WAKO) under ice cooling, warmed to room temperature, stirred and for 19.5 hours, then warmed to 40°C, and further stirred for 4 hours. The reaction mixture was added dropwise with a solution of 2-amino-6-bromonaphthalene (443 mg) in benzene (5 ml) under ice cooling, warmed to

room temperature, and stirred 21.5 hours. The reaction mixture was filtered to obtain solid, and the solid was added with ethyl acetate, mixed and filtered again. The solvent of the filtrate was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 2:1) to obtain the title compound (Intermediate 66, 158 mg).

Synthesis of methyl 3-[4-cyclopentylmethoxy-3-(6-sulfamoylaminonaphthalen-2-yl)phenyl]propionate (Compound No. V-22)

According to a procedure described in literature (Miyaura, N. et al., Tetrahedron.Lett., 1997, p.3447), Compound No. A-1 (209 mg), bis(pinacolate)diboron (177 mg, Ald), [1,1'-bis(diphenylphosphono)ferrocene]palladium(II) dichloride (hereinafter abbreviated as "PdCl<sub>2</sub>(dppf)", 28 mg, TCI) and potassium acetate (182.3 mg, Ald) were added to DMF (6 ml), and heated to 80°C with stirring under argon gas atmosphere for 5 hours. The reaction mixture was cooled to room temperature, then added with Intermediate 91 (130 mg), PdCl<sub>2</sub>(dppf) (30 mg) and 2 M aqueous sodium carbonate (0.9 ml), and heated to 80°C for 21 hours with stirring under argon gas atmosphere. The reaction mixture was added with ethyl acetate (100 ml), washed with saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 3:1) to obtain the title compound (Compound No. V-22, 46 mg).

[Example V-23]

Synthesis of 3-[4-cyclopentylmethoxy-3-(6-sulfamoylaminonaphthalen-2-yl)phenyl]propionic acid (Compound No. V-23)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 24 hours, Compound No. V-22 (41 mg) and 2 N aqueous sodium hydroxide (340  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. V-23, 22 mg).

[Example V-27]

Synthesis of methyl 3-[4-cyclopentyloxy-3-(1H-indol-5-yl)phenyl]propionate  
(Compound No. V-27)

According to the procedure described in the synthesis method of Compound No. C-1 with the modifications that the reaction was carried out at 80°C for 5 hours, and the purification was performed by flash column chromatography (hexane:ethyl acetate = 5:1), Compound No. A-5 (367 mg), 5-indoleboronic acid (310 mg, Frontier), 2 M aqueous sodium carbonate (0.9 ml) and (Ph<sub>3</sub>P)<sub>4</sub>Pd (132 mg) were reacted and treated to obtain the title compound (Compound No. V-27, 340 mg).

[Example V-28]

Synthesis of 3-[4-cyclopentyloxy-3-(1H-indol-5-yl)phenyl]propionic acid (Compound No. V-28)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 2 hours, Compound No. V-27 (330 mg) and 2 N aqueous sodium hydroxide (1.40 ml) were reacted and treated to obtain the title compound (Compound No. V-28, 310 mg).

[Example V-29]

Synthesis of methyl 3-[4-cyclopentyloxy-3-(1-methyl-1H-indol-5-yl)phenyl]propionate (Compound No. V-29)

A solution of Compound No. V-27 (123 mg) in DMF (5 ml) was added with 60% sodium hydride (19 mg) under ice cooling, and stirred for 10 minutes. The reaction mixture was added dropwise with methyl iodide (100  $\mu$ l), stirred for 10 minutes, then warmed to room temperature, and further stirred for 1 hour. The reaction mixture was poured into ice water, and ethyl acetate (100 ml) was added for extraction. The organic layer was successively washed with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride, and saturated brine and dried, and then the solvent was evaporated under reduced pressure.



The residue was purified by flash column chromatography (hexane:ethyl acetate = 8:1) to obtain the title compound (Compound No. V-29, 126 mg).

[Example V-30]

Synthesis of 3-[4-cyclopentyloxy-3-(1-methyl-1H-indol-5-yl)phenyl]propionic acid (Compound No. V-30)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 1 hour, Compound No. V-29 (123 mg) and 2 N aqueous sodium hydroxide (330  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. V-30, 110 mg).

[Example V-31]

Synthesis of methyl 3-[4-cyclopentylmethoxy-3-(1H-indol-4-yl)phenyl]propionate (Compound No. V-31)

According to the procedure described in the synthesis method of Compound No. C-1 with the modifications that the reaction was carried out for 21 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 6:1), Compound No. A-1 (200 mg), 4-indoleboronic acid (170 mg) obtainable from 4-bromoindole (TCI) according to a known method described in a publication (Doll, M. et al., J. Org. Chem, 1999, vol. 64, p.1372), 2 M aqueous sodium carbonate (550  $\mu$ l) and (Ph<sub>3</sub>P)<sub>4</sub>Pd (60 mg) were reacted and treated to obtain the title compound (Compound No. V-31, 214 mg).

[Example V-32]

Synthesis of 3-[4-cyclopentylmethoxy-3-(1H-indol-4-yl)phenyl]propionic acid (Compound No. V-32)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 1 hour, Compound No. V-31 (210 mg) and 2 N aqueous sodium hydroxide (0.60 ml) were reacted and treated to obtain the title compound (Compound No. V-32, 173 mg).

[Example V-33]

Synthesis of 4-bromo-1-methyl-1H-indole (Intermediate 67)

According to the procedure described in the synthesis method of Compound No. V-29 with the modifications that the reaction was carried out for 30 minutes, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 10:1), 4-bromoindole (5 g), 60% sodium hydride (1.14 g) and methyl iodide (3.18 ml, TCI) were reacted and treated to obtain the title compound (Intermediate 67, 4.95 g).

Synthesis of 1-methyl-1H-indole-4-boronic acid (Intermediate 68)

A solution of Intermediate 67 (4.90 g) in anhydrous THF (30 ml) was cooled to -78°C under argon gas atmosphere, then added dropwise with a 1.62 M solution of t-butyllithium in pentane (28.8 ml) over 30 minutes, and stirred for 30 minutes. This reaction mixture was added dropwise with (iPrO)<sub>3</sub>B (10.77 ml) over 10 minutes, stirred for 1 hour, then warmed to room temperature, and further stirred for 2.5 hours. The reaction mixture was poured into 1.2 N aqueous phosphoric acid (250 ml) containing ice, and extracted with diethyl ether (200 ml x 3). The organic layer was extracted with 0.4 N aqueous sodium hydroxide (150 ml x 3), and the aqueous layer was made acidic with 5 N hydrochloric acid under ice cooling, and extracted with diethyl ether (200 ml x 3) again. The organic layer was washed with saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was washed with hexane to obtain the title compound (Intermediate 68, 3.17 g).

Synthesis of methyl 3-[4-cyclopentylmethoxy-3-(1-methyl-1H-indol-4-yl)phenyl]propionate (Compound No. V-33)

According to the procedure described in the synthesis method of Compound No. C-1 with the modifications that the reaction was carried out for 18 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl

acetate = 9:1), Compound No. A-1 (200 mg), Intermediate 68 (185 mg), 2 M aqueous sodium carbonate (550  $\mu$ l) and (Ph<sub>3</sub>P)<sub>4</sub>Pd (60 mg) were reacted and treated to obtain the title compound (Compound No. V-33, 208 mg).

[Example V-34]

Synthesis of 3-[4-cyclopentylmethoxy-3-(1-methyl-1H-indol-4-yl)phenyl]propionic acid (Compound No. V-34)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 3 hours, Compound No. V-33 (200 mg) and 2 N aqueous sodium hydroxide (0.60 ml) were reacted and treated to obtain the title compound (Compound No. V-34, 182 mg).

[Example V-43]

Synthesis of 3-[4-cyclopentylmethoxy-3-[1-(2-hydroxyethyl)-1H-indol-5-yl]phenyl]propionic acid (Compound No. V-43)

According to the procedure described in the synthesis method of Compound No. V-29 with the modifications that the reaction was carried out for 1.5 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 8:1), Compound No. V-27 (144mg), 60% sodium hydride (38 mg) and ethyl bromoacetate (160  $\mu$ l, TCI) were reacted and treated to obtain an oily substance. This substance was reacted with 2 N aqueous sodium hydroxide (300  $\mu$ l) and treated according to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 1 hour to obtain the title compound (Compound No. V-43, 36 mg).

[Example V-44]

Synthesis of methyl 3-[4-cyclopentylmethoxy-3-(3-formyl-1H-indol-5-yl)phenyl]propionate (Compound No. V-44)

A solution of Compound No. V-27 (75 mg) in DMF (6 ml) was added dropwise with phosphoryl chloride (30  $\mu$ l, TCI) under ice cooling, stirring for 1

hour, then warmed to 35°C, and further stirred for 1 hour. The reaction mixture was added with 1 N aqueous sodium hydroxide (3 ml) containing ice, and extracted with ethyl acetate (90 ml). The organic layer was washed with saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography (hexane:ethyl acetate = 5:1) to obtain the title compound (Compound No. V-44, 86 mg).

[Example V-45]

Synthesis of 3-[4-cyclopentylmethoxy-3-(3-formyl-1H-indol-5-yl)phenyl]propionic acid (Compound No. V-45)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 3 hours, Compound No. V-44 (86 mg) and 2 N aqueous sodium hydroxide (110  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. V-45, 60 mg).

[Example V-47]

Synthesis of methyl 3-[3-(3-acetyl-1H-indol-5-yl)-4-cyclopentylmethoxyphenyl]propionate (Compound No. V-47)

A solution of Compound No. V-27 (98 mg) in methylene chloride (2 ml) was added with aluminum chloride (81 mg, Ald) and acetyl chloride (60  $\mu$ l), and stirred for 4 hours. The reaction mixture was added with 1 N hydrochloric acid (2 ml), and extracted with methylene chloride (60 ml). The organic layer was washed with saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography (hexane:ethyl acetate = 4:1) to obtain the title compound (Compound No. V-47, 47 mg).

[Example V-48]

Synthesis of 3-[3-(3-acetyl-1H-indol-5-yl)-4-cyclopentylmethoxyphenyl]propionic acid (Compound No. V-48)

According to the procedure described in the synthesis method of

Intermediate 9 provided that the reaction was carried out for 4 hours, Compound No. V-47 (45 mg) and 2 N aqueous sodium hydroxide (110  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. V-48, 44 mg).

[Example V-50]

Synthesis of methyl 3-[4-cyclopentylmethoxy-3-(3-methyl-1H-indol-5-yl)phenyl]propionate (Compound No. V-50)

According to the procedure described in the synthesis method of Intermediate 95 with the modifications that the reaction was carried out for 13 hours, and the purification was performed by flash column chromatography (hexane:ethyl acetate = 4:1), 5-bromo-3-methylindole (1.63 g) obtainable from 5-bromoindole (TCI) by a known method described in a publication (Wayland, E.N., J. Org. Chem, 1967, vol. 32, p.828) was reacted with 30% potassium hydride (1.08 g), a 1.7 M solution of t-butyllithium in pentane (9.7 ml) and (*i*PrO)<sub>3</sub>B (3.75 ml) and treated to obtain crude 3-methyl-5-indoleboronic acid. This compound was reacted with Compound No. A-1 (803 mg), 2 M aqueous sodium carbonate (2 ml) and (Ph<sub>3</sub>P)<sub>4</sub>Pd (241 mg) and treated to obtain the title compound (Compound No. V-50, 552 mg).

[Example V-51]

Synthesis of 3-[4-cyclopentylmethoxy-3-(3-methyl-1H-indol-5-yl)phenyl]propionic acid (Compound No. V-51)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 2 hours, Compound No. V-50 (130 mg) and 2 N aqueous sodium hydroxide (370  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. V-51, 127 mg).

[Example V-54]

Synthesis of 4-bromo-1H-indazole (Intermediate 69)

According to a known method described in a publication (Schumann, P. et

al., Bioorg. Med. Chem. Lett., 2001, vol. 11, p.1153), the title compound (Intermediate 69, 1.68 g) was obtained from commercially available 3-bromotoluidine (4.51 g, Ald).

Synthesis of methyl 3-[4-cyclopentyloxy-3-(1H-indazol-4-yl)phenyl]propionate (Compound No. V-54)

According to the procedure described in the synthesis method of Compound No. V-22 provided that the purification was performed by flash column chromatography (hexane:ethyl acetate = 2:1), Compound No. A-5 (328 mg), bis(pinacolate)diboron (281 mg), PdCl<sub>2</sub>(dppf) (61 mg) and potassium acetate (303 mg) were reacted at 80°C for 4 hours, and then this reaction mixture was added with Intermediate 105 (161 mg), PdCl<sub>2</sub>(dppf) (64 mg) and 2 M aqueous sodium carbonate (1.5 ml), reacted at 80°C for 9 hours and treated to obtain the title compound (Compound No. V-54, 111 mg).

[Example V-55]

Synthesis of 3-[4-cyclopentyloxy-3-(1H-indazol-4-yl)phenyl]propionic acid (Compound No. V-55)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 2 hours, Compound No. V-54 (108 mg) and 2 N aqueous sodium hydroxide (400  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. V-55, 99 mg).

[Example V-57]

Synthesis of 4-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)-2-methylnitrobenzene (Intermediate 70)

According to the procedure described in the synthesis method of Compound No. V-22, 5-bromo-2-nitrotoluene (4.30 g) synthesized by nitrating 3-bromotoluene (WAKO) by a known method, bis(pinacolate)diboron (5.59 g), PdCl<sub>2</sub>(dppf) (440 mg) and potassium acetate (6.09 g) were heated with stirring at 80°C for 3 hours under

argon gas atmosphere. The reaction mixture was added with ethyl acetate (300 ml), and washed successively with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride and saturated brine. The organic layer was dried, and then the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography (hexane:ethyl acetate = 8:1) to obtain the title compound (Intermediate 70, 4.21 g).

Synthesis of 4-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)-2-methylaniline  
(Intermediate 71)

According to the procedure described in the synthesis method of Compound No. Q-2 with the modification that the reaction was carried out for 30 minutes, Intermediate 70 (4.20 g) and platinum oxide (50 mg) were added, then reacted and treated under hydrogen atmosphere to obtain the title compound (Intermediate 71, 2.81 g).

Synthesis of methyl 3-(4'-amino-6-cyclopentyloxy-3'-methlbiphenyl-3-yl)propionate  
(Intermediate 72)

According to the procedure described in the synthesis method of Compound No. C-1 with the modifications that the reaction was carried out for 15.5 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 6:1), Compound No. A-5 (701mg), Intermediate 71 (604mg), 2 M aqueous sodium carbonate (1.8ml), and (Ph<sub>3</sub>P)<sub>4</sub>Pd (182mg) were reacted and treated to obtain the title compound (Intermediate 72, 762 mg).

Synthesis of methyl 3-[4-cyclopentyloxy-3-(1H-indazol-5-yl)phenyl]propionate  
(Compound No. V-57)

A solution of Intermediate 72 (760 mg) in acetic acid (4 ml) was added with an aqueous solution (0.7 ml) of sodium nitrite (156 mg) under ice cooling, and stirred for 30 minutes. This reaction mixture was added with urea (350 mg), warmed to room temperature, stirred for 30 minutes, then added with toluene (8

ml) and water (4 ml), and further stirred for 60 hours. The reaction mixture was extracted with toluene (50 ml x 2). The organic layer was washed successively with saturated aqueous sodium hydrogencarbonate, and saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 6:1) to obtain the title compound (Compound No. V-57, 411 mg).

[Example V-58]

Synthesis of 3-[4-cyclopentyloxy-3-(1H-indazol-5-yl)phenyl]propionic acid  
(Compound No. V-58)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 2.5 hours, Compound No. V-57 (86 mg) and 2 N aqueous sodium hydroxide (250  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. V-58, 82 mg).

[Example V-66]

Synthesis of 5-bromo-3-methyl-1H-indazole (Intermediate 73)

According to the procedure described in the synthesis method of Compound No. V-57 provided that the reaction was carried out for 121 hours, 4-bromo-2-ethylaniline (5.01 g, LANC) and sodium nitrite (1.918 g) were reacted and treated to obtain the title compound (Intermediate 73, 3.30 g).

Synthesis of methyl 3-[4-cyclopentyloxy-3-(3-methyl-1H-indazol-5-yl)phenyl]propionate (Compound No. V-66)

According to the procedure described in the synthesis method of Compound No. V-22 provided that the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 5:2), Compound No. A-5 (434 mg), bis(pinacolate)diboron (367 mg), PdCl<sub>2</sub>(dppf) (101 mg), and potassium acetate (339 mg) were reacted at 80°C for 4 hours, and then this reaction mixture was added with Intermediate 108 (273 mg), PdCl<sub>2</sub>(dppf) (104 mg) and 2 M aqueous sodium



carbonate (1.1 ml), reacted at 80°C for 18 hours and treated to obtain the title compound (Compound No. V-66, 98 mg).

[Example V-67]

Synthesis of 3-[4-cyclopentyloxy-3-(3-methyl-1H-indazol-5-yl)phenyl]propionic acid (Compound No. V-67)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 2 hours, Compound No. V-66 (97 mg) and 2 N aqueous sodium hydroxide (400  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. V-67, 54 mg).

[Example V-68]

Synthesis of methyl 3-[4-cyclopentyloxy-3-(1,3-dimethyl-1H-indazol-5-yl)phenyl]propionate (Compound No. V-68)

According to the procedure described in the synthesis method of Compound No. V-29 with the modifications that the reaction was carried out for 16 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 3:1), Compound No. V-66 (112 mg), 60% sodium hydride (24 mg) and methyl iodide (95  $\mu$ l) were reacted and treated to obtain the title compound (Intermediate 110, 45 mg).

[Example V-69]

Synthesis of 3-[4-cyclopentyloxy-3-(1,3-dimethyl-1H-indazol-5-yl)phenyl]propionic acid (Compound No. V-69)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 3 hours, Compound No. V-68 (45 mg) and 2 N aqueous sodium hydroxide (120  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. V-69, 42 mg).

[Example V-73]

Synthesis of methyl 3-[3-(benzo[b]thiophen-5-yl)-4-

cyclopentylmethyloxyphenyl]propionate (Compound No. V-73)

According to the procedure described in the synthesis method of Compound No. V-22 provided that the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 10:1), Compound No. A-1 (371 mg), bis(pinacolate)diboron (294 mg), PdCl<sub>2</sub>(dppf) (67 mg) and potassium acetate (308 mg) were reacted at 80°C for 10 hours, and then this reaction mixture was added with 5-bromobenzo[b]thiophene (301.4 mg) obtainable from 4-bromothiophenol (TCI) by a known method described in a publication (Seed, A.J., J. Mater. Chem., 2000, vol. 10, p.2069), PdCl<sub>2</sub>(dppf) (65 mg) and 2 M aqueous sodium carbonate (0.9 ml), reacted at 80°C for 16 hours and treated to obtain the title compound (Compound No. V-73, 97 mg).

[Example V-74]

Synthesis of 3-[3-(benzo[b]thiophen-5-yl)-4-cyclopentylmethyloxyphenyl]propionic acid (Compound No. V-74)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 3 hours, Compound No. V-73 (95 mg) and 2 N aqueous sodium hydroxide (250  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. V-74, 93 mg).

[Example V-77]

Synthesis of (3-bromophenyl)thiourea (Intermediate 74)

A solution of 3-bromoaniline (10.89 ml, TCI) in 20% aqueous hydrochloric acid (18.2 ml) was added with ammonium thiocyanate (8.02 g, WAKO) and sodium hydrogensulfite (701 mg, WAKO), and stirred at 100°C for 22 hours. The reaction mixture was added with chloroform (20 ml) for extraction, and the organic layer was dried. Then, the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 2:1) to obtain the title compound (Intermediate 74, 4.45 g).

#### Synthesis of 2-amino-5-bromobenzothiazole (Intermediate 75)

A solution of Intermediate 74 (1.29 g) in chloroform (12 ml) was added dropwise with a solution of bromine (272  $\mu$ l, WAKO) in chloroform (1.5 ml), refluxed by heating for 2.5 hours, and stirred at room temperature for 16 hours. The reaction mixture was concentrated under reduced pressure, neutralized with 5% aqueous ammonia, and then added with water (50 ml) and methylene chloride (150 ml) for extraction. The organic layer was dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 2:1) to obtain the title compound (Intermediate 75, 609 mg).

#### Synthesis of methyl 3-[3-(2-aminobenzothiazol-5-yl)-4-cyclopentylmethoxyphenyl]propionate (Compound No. V-77)

A solution of Intermediate 75 (459.1 mg) in anhydrous THF (30 ml) was added with N,N,N',N'-tetramethylethylenediamine (1.51 ml, WAKO), cooled to -78°C under argon gas atmosphere, then added dropwise with a 1.62 M solution of t-butyllithium in pentane (7.06 ml), and stirred for 30 minutes. The reaction mixture was added dropwise with (iPrO)<sub>3</sub>B (2.77 ml), stirred for 30 minutes, then warmed to room temperature, and further stirred for 1.5 hours. The reaction mixture was added with 0.5 M aqueous sulfuric acid (7.5 ml) and extracted with diethyl ether (50 ml x 3). The organic layer was washed with saturated brine and dried, and then the solvent was evaporated under reduced pressure to obtain crude 2-amino-5-benzothiazoleboronic acid. This compound was reacted with Compound No. A-1 (344 mg), 2 M aqueous sodium carbonate (4.5 ml) and (Ph<sub>3</sub>P)<sub>4</sub>Pd (179 mg) and treated according to the procedure described in the synthesis method of Compound No. V-3 with the modifications that the reaction was carried out for 12 hours, and the purification was performed by flash column chromatography (hexane:ethyl acetate = 2:1) to obtain the title compound (Compound No. V-77, 76

mg).

[Example V-78]

Synthesis of 3-[3-(2-aminobenzothiazol-5-yl)-4-cyclopentylmethoxyphenyl]propionic acid (Compound No. V-78)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 2.5 hours, Compound No. V-77 (77 mg) and 2 N aqueous sodium hydroxide (380  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. V-78, 69 mg).

[Example V-79]

Synthesis of ethyl 3-[3-(benzothiazol-5-yl)-4-cyclopentylmethoxyphenyl]propionate (Compound No. V-79)

A solution of Compound No. V-77 (215 mg) in acetonitrile (10 ml) was added with 30% aqueous hypophosphorous acid (3 ml, WAKO), cooled to 0°C, added dropwise with an aqueous solution (1 ml) of sodium nitrite (187 mg), stirred for 30 minutes, then warmed to room temperature, and further stirred for 20 hours. The reaction mixture was poured into water (50 ml), neutralized by addition of 2 N aqueous sodium hydroxide, and added with ethyl acetate (90 ml x 3) for extraction. The organic layer was washed with saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 10:1) to obtain the title compound (Compound No. V-79, 78 mg).

[Example V-80]

Synthesis of 3-[3-(benzothiazol-5-yl)-4-cyclopentylmethoxyphenyl]propionic acid (Compound No. V-80)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 2 hours, Compound No. V-79 (75 mg) and 2 N aqueous sodium hydroxide (500  $\mu$ l) were reacted and

treated to obtain the title compound (Compound No. V-80, 66 mg).

[Example V-81]

Synthesis of methyl 3-[4-cyclopentylmethoxy-3-(2-methylbenzothiazol-5-yl)phenyl]propionate (Compound No. V-81)

According to the procedure described in the synthesis method of Compound No. V-13 with the modifications that the reaction was carried out for 13 hours, and the purification was performed by flash column chromatography (hexane:ethyl acetate = 5:1), crude 2-methyl-5-benzothiazoleboronic acid prepared from 5-bromo-2-methylbenzothiazole (684 mg, TCI), a 1.7 M solution of t-butyllithium in pentane (7.06 ml) and  $(i\text{-PrO})_3\text{B}$  (3.46 ml), Compound No. A-1 (515 mg), 2 M aqueous sodium carbonate (6.5 ml) and  $(\text{Ph}_3\text{P})_4\text{Pd}$  (258 mg) were reacted and treated to obtain the title compound (Compound No. V-81, 240 mg).

[Example V-82]

Synthesis of 3-[4-cyclopentylmethoxy-3-(2-methylbenzothiazol-5-yl)phenyl]propionic acid (Compound No. V-82)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 4 hours, Compound No. V-81 (227 mg) and 2 N aqueous sodium hydroxide (1.11 ml) were reacted and treated to obtain the title compound (Compound No. V-82, 132 mg).

[Example V-83]

Synthesis of ethyl 3-[4-cyclopentylmethoxy-3-[2-(N,N-dimethylamino)benzothiazol-6-yl]phenyl]propionate (Compound No. V-83)

According to the procedure described in the synthesis method of Compound No. V-29 with the modifications that the reaction was carried out for 4 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 7:1), Compound No. V-77 (155 mg), 60% sodium hydride (16 mg) and methyl iodide (68.5  $\mu\text{l}$ ) were reacted and treated to obtain the title compound

(Compound No. V-83, 48 mg).

[Example V-84]

Synthesis of 3-[4-cyclopentylmethyloxy-3-[2-(N,N-dimethylamino)benzothiazol-6-yl]phenyl]propionic acid (Compound No. V-84)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 3 hours, Compound No. V-83 (47 mg) and 2 N aqueous sodium hydroxide (200  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. V-84, 35 mg).

[Example V-88]

Synthesis of ethyl 3-[3-(2-bromobenzothiazol-6-yl)-4-cyclohexylmethyloxyphenyl]propionate (Intermediate 76)

A solution obtained beforehand by adding t-butyl nitrite (178  $\mu$ l, TCI) and copper(I) bromide (241 mg, WAKO) to acetonitrile (10 ml) and mixing them was added dropwise with a solution of Compound No. V-83 (381 mg) in acetonitrile (5 ml) and stirred at room temperature for 1.5 hours. The solvent of the reaction mixture was concentrated under reduced pressure, and the residue was purified by column chromatography (Quad, hexane:ethyl acetate = 10:1) to obtain the title compound (Intermediate 76, 341 mg).

Synthesis of 3-[4-cyclopentylmethyloxy-3-(2-methoxybenzothiazol-6-yl)phenyl]propionic acid (Compound No. V-88)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 18 hours, Intermediate 76 (169 mg) and 2 N aqueous sodium hydroxide (500  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. V-88, 114 mg).

[Example V-89]

Synthesis of 3-[4-cyclopentylmethyloxy-3-(2-oxo-2,3-dihydrobenzothiazol-6-yl)phenyl]propionic acid (Compound No. V-64)

A solution of Intermediate 76 (202 mg) in ethanol (8 ml) was added with 5 N aqueous hydrochloric acid (1.5 ml), and stirred at 80°C for 18.5 hours. The reaction mixture was concentrated under reduced pressure, and added with water (20 ml) and ethyl acetate (80 ml) for extraction. The organic layer was washed with saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was added with 2 N aqueous sodium hydroxide (1.0 ml), reacted and treated according to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 2 hours to obtain the title compound (Compound No. V-89, 250 mg).

[Example V-91]

Synthesis of 3-[4-cyclopentylmethoxy-3-(2-thioxo-2,3-dihydrobenzothiazol-6-yl)phenyl]propionic acid (Compound No. V-91)

A solution obtained beforehand by adding thiourea (52 mg, WAKO) to 1 M sulfuric acid (5 ml) and mixing them was added with a solution of Intermediate 76 (101 mg) in acetonitrile (5 ml), and stirred at 90°C for 20 hours. The reaction mixture was poured into water (20 ml), neutralized by addition of 1 N aqueous sodium hydroxide under ice cooling, and then extracted with ethyl acetate (80 ml x 3). The organic layer was washed with saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, methylene chloride:ethanol = 30:1) to obtain the title compound (Compound No. V-91, 46 mg).

Synthesis examples for compounds used for preparation of the compounds mentioned in the examples are shown below.

Syntheses of 4-bromo-1-methyl-1H-indazole (Intermediate 77) and 4-bromo-2-methyl-2H-indazole (Intermediate 78)

According to the procedure described in the synthesis method of Compound No. V-29 with the modifications that the reaction was carried out for 8 hours, and

the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 5:1), Intermediate 69 (600 mg), 60% sodium hydride (191 mg), and methyl iodide (379  $\mu$ l) were reacted and treated to obtain the title compounds (Intermediate 119, 432mg and Intermediate 120, 164 mg).

#### Synthesis of 5-bromo-1H-indazole (Intermediate 79)

The title compound (Intermediate 121, 0.91 g) was obtained from commercially available 4-bromotoluidine (3.33 g, Ald) by a method known from the aforementioned literature (Bioorg. Med. Chem. Lett., 2001, vol. 11, p.1153).

#### Syntheses of 5-bromo-1-methyl-1H-indazole (Intermediate 80) and 5-bromo-2-methyl-2H-indazole (Intermediate 81)

According to the procedure described in the synthesis method of Compound No. V-29 with the modifications that the reaction was carried out for 4.5 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 5:1), Intermediate 79 (300 mg), 60% sodium hydride (80 mg), and methyl iodide (161  $\mu$ l) were reacted and treated to obtain the title compounds (Intermediate 80, 201mg and Intermediate 81, 87 mg).

#### Synthesis of 1-methyl-1H-indazole-5-boronic acid (Intermediate 82)

According to the procedure described in the synthesis method of Compound No. V-3, Intermediate 80 (1.69 g), a 1.6 M solution of n-butyllithium in hexane (7.50 ml) and (iPrO)<sub>3</sub>B (3.23 ml) were reacted and treated to obtain the title compound (Intermediate 82, 1.39 g).

#### Syntheses of 5-bromo-1-ethyl-1H-indazole (Intermediate 83) and 5-bromo-2-ethyl-2H-indazole (Intermediate 84)

According to the procedure described in the synthesis method of Compound No. V-29 with the modifications that the reaction was carried out for 2 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 5:1), Intermediate 79 (420 mg), 60% sodium hydride (111 mg), and ethyl



iodide (375  $\mu$ l) were reacted and treated to obtain the title compounds (Intermediate 83, 250mg and Intermediate 84, 127 mg).

#### Synthesis of 6-bromo-1H-indazole (Intermediate 85)

The title compound was obtained from commercially available 5-bromotoluidine (3.33 g, Ald) by the method known from the aforementioned literature (Bioorg. Med. Chem. Lett., 2001, vol. 11, p.1153) to obtain the title compound (Intermediate 85, 0.42 g).

#### Syntheses of 6-bromo-1-methyl-1H-indazole (Intermediate 86) and 6-bromo-2-methyl-2H-indazole (Intermediate 87)

According to the procedure described in the synthesis method of Compound No. V-29 with the modifications that the reaction was carried out for 2.5 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 5:1), Intermediate 85 (277 mg), 60% sodium hydride (86 mg), and methyl iodide (175  $\mu$ l) were reacted and treated to obtain the title compounds (Intermediate 86, 196 mg and Intermediate 87, 89 mg).

#### Synthesis of 5-bromo-2-t-butylthiobenzaldehyde (Intermediate 88)

A solution of 5-bromo-2-fluorobenzaldehyde (4.06 g, Avocado) in 2-propanol (20 ml) was added with 2-methyl-2-propanethiol (2.26 ml, Ald) and potassium carbonate (3.04 g), and heated with stirring for 18 hours. The reaction mixture was cooled to room temperature, then poured into water (50 ml), and extracted with chloroform (75 ml x 3). The organic layer was washed twice with saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography (hexane:ethyl acetate = 20:1) to obtain the title compound (Intermediate 88, 754 mg).

#### Synthesis of 5-bromobenzo[d]isothiazole (Intermediate 89)

A solution obtained beforehand by mixing 2 N aqueous sodium hydroxide (2.19 ml) in an aqueous solution (5 ml) of hydroxylamine hydrochloride (308 mg,

WAKO) was added dropwise to a solution of Intermediate 88 (401 mg) in ethanol (5 ml) at room temperature over 15 minutes. The reaction mixture was refluxed by heating for further 2 hours, then cooled to room temperature, poured into water (30 ml), and extracted with ethyl acetate (100 ml x 3). The organic layer was washed successively with aqueous saturated ammonium chloride aqueous, saturated aqueous sodium hydrogencarbonate, and saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was added with polyphosphoric acid (21.4 g), and heated with stirring at 100°C for 2 hours. The reaction mixture was poured into ice water (100 ml), neutralized with 5 N aqueous sodium hydroxide under ice cooling, and then extracted with ethyl acetate (100 ml x 3). The organic layer was washed twice with saturated brine and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography (hexane:ethyl acetate = 20:1) to obtain the title compound (Intermediate 89, 143 mg).

#### Synthesis of 5-bromobenzo[c]isothiazole (Intermediate 90)

A solution of methanesulfonamide (5.34 g, TCI) in dehydrated benzene (9 ml) was added with thionyl chloride (6.0 ml) under ice cooling, and refluxed by heating for 24 hours. The reaction mixture was concentrated under reduced pressure, and a solution of the residue in dehydrated benzene (4 ml) was added dropwise to a solution of 4-bromotoluidine (1.49 g) in dehydrated benzene (40 ml) under ice cooling. This mixture was added dropwise with a solution of pyridine (0.97 ml) in dehydrated benzene (4 ml) under ice cooling, and refluxed by heating for 80 hours under argon gas atmosphere. The reaction mixture was cooled to room temperature, poured into water (100 ml), and extracted with chloroform (100 ml x 3). The organic layer was dried, and then the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography (hexane:ethyl acetate = 10:1) to obtain the title compound (Intermediate 90, 618

mg).

#### Synthesis of 6-bromoimidazo[1,2-a]pyridine (Intermediate 91)

The title compound (Intermediate 91, 3.36 g) was obtained from commercially available bromoacetaldehyde-diethylacetal (4.7 ml, WAKO) and 2-amino-5-bromopyridine (4.32 g, Ald) by a known method described in a publication (Yamanaka, M. et al., Chem. Pharm. Bull., 1991, vol. 39, p.1556).

#### Synthesis of 5-bromo-1H-pyrrolo[2,3-b]pyridine (Intermediate 92)

The title compound (Intermediate 92, 182 mg) was obtained from commercially available 1H-pyrrolo[2,3-b]pyridine (1.3 g, TCI) by a known method described in a publication (Mazeas, D. et al, Heterocycles, 1999, vol. 50, p.1065).

#### Synthesis of 5-bromo-1-methyl-1H-pyrrolo[2,3-b]pyridine (Intermediate 93)

According to the procedure described in the synthesis method of Compound No. V-29 with the modifications that the reaction was carried out for 2 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 15:1), Intermediate 92 (98 mg), 60% sodium hydride (33 mg), and methyl iodide (53  $\mu$ l) were reacted and treated to obtain the title compound (Intermediate 93, 88 mg).

#### Synthesis of 6-bromoisoquinoline (Intermediate 94)

The title compound (Intermediate 94, 1.46 g) was obtained from commercially available 4-bromobenzaldehyde (15.0 g, WAKO) by a known method described in a publication (Nerenz, H. et al., J. Chem. Soc. Perkin Trans. 2, 1998, p.437).

#### Synthesis of 6-bromo-2H-isoquinolin-1-one (Intermediate 95)

A solution of Intermediate 94 (1.04 g) in methylene chloride (3 ml) was added with a solution of 3-chloroperbenzoic acid (2.16 g) in methylene chloride (3 ml), and stirred for 20 hours. The reaction mixture was added with methylene chloride (200 ml), and washed successively with saturated aqueous sodium

hydrogencarbonate, saturated aqueous ammonium chloride and saturated brine. The organic layer was dried, and then the solvent was evaporated under reduced pressure. A solution of the residue in acetic anhydride (10 ml) was refluxed by heating for 5 hours. The reaction mixture was concentrated under reduced pressure, and then the residue was added with 2.5 N aqueous sodium hydroxide (20 ml), and stirred at 100°C for 1 hour. The reaction mixture was cooled to room temperature, and neutralized with 5 N aqueous hydrochloric acid under ice cooling to obtain the precipitated title compound (Intermediate 95, 623 mg).

[Examples V-1 to V-115]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification including the examples described above are shown in Table-V-1 to Table-V-3.

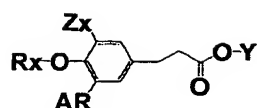


Table-V-1

Exp.	RxO	Y	Zx	AR	Syn	LCMS		
						method	RTime	Mass
V-1	cHexMeO	Et	H	1-Nap	V-1			
V-2	cHexMeO	H	H	1-Nap	V-2	C		375 (M <sup>+</sup> +1)
V-3	cPenMeO	Me	H	6OH-2-Nap	V-3			
V-4	cPenMeO	H	H	6OH-2-Nap	V-4			
V-5	cPenMeO	Me	H	5OH-2-Nap	V-5			
V-6	cPenMeO	H	H	5OH-2-Nap	V-6			
V-7	cPenMeO	Me	H	7OH-2-Nap	V-7			
V-8	cPenMeO	H	H	7OH-2-Nap	V-8			
V-9	cPenMeO	Me	H	6OMe-2-Nap	V-1			
V-10	cPenMeO	H	H	6OMe-2-Nap	V-2	C		418(M <sup>+</sup> )
V-11	cPenMeO	Me	H	6(OCH <sub>2</sub> CONMe <sub>2</sub> )-2-Nap	V-11			
V-12	cPenMeO	H	H	6(OCH <sub>2</sub> CONMe <sub>2</sub> )-2-Nap	V-12			
V-13	cPenMeO	Me	H	6NH <sub>2</sub> -2-Nap	V-13			
V-14	cPenMeO	H	H	6NH <sub>2</sub> -2-Nap	V-14			
V-15	cPenMeO	H	H	6(NMe <sub>2</sub> )-2-Nap	V-13,V-14	C		418(M <sup>+</sup> +1)
V-16	cPenMeO	H	H	6(NHCOCH <sub>2</sub> OH)-2-Nap	V-16			
V-17	cPenMeO	H	H	6(NHCO-2-Furan)-2-Nap	V-16	C		484(M <sup>+</sup> +1)
V-18	cPenMeO	Me	H	6(NHSO <sub>2</sub> Me)-2-Nap	V-18			
V-19	cPenMeO	H	H	6(NHSO <sub>2</sub> Me)-2-Nap	V-19			
V-20	cPenMeO	Me	H	6(NHSO <sub>2</sub> NMe <sub>2</sub> )-2-Nap	V-20			
V-21	cPenMeO	H	H	6(NHSO <sub>2</sub> NMe <sub>2</sub> )-2-Nap	V-21			
V-22	cPenMeO	Me	H	6(NHSO <sub>2</sub> NH <sub>2</sub> )-2-Nap	V-22			
V-23	cPenMeO	H	H	6(NHSO <sub>2</sub> NH <sub>2</sub> )-2-Nap	V-23			
V-24	cPenMeO	H	H	6(SO <sub>2</sub> Me)-2-Nap	V-22,V-23	C		452(M <sup>+</sup> )
V-25	cPenMeO	H	H	6(SO <sub>2</sub> NH <sub>2</sub> )-2-Nap	V-22,V-23	C		453(M <sup>+</sup> )
V-26	cPenMeO	H	H	6(SO <sub>2</sub> NHMe)-2-Nap	V-22,V-23	C		468(M <sup>+</sup> +1)
V-27	cPenO	Me	H	5-Ind	V-27			
V-28	cPenO	H	H	5-Ind	V-28			
V-29	cPenO	Me	H	1Me-5-Ind	V-29			
V-30	cPenO	H	H	1Me-5-Ind	V-30			
V-31	cPenMeO	Me	H	4-Ind	V-31			
V-32	cPenMeO	H	H	4-Ind	V-32			
V-33	cPenMeO	Me	H	1Me-4-Ind	V-33			
V-34	cPenMeO	H	H	1Me-4-Ind	V-34			
V-35	cPenMeO	H	H	6-Ind	V-31,V-32	C		377(M <sup>+</sup> )
V-36	cPenMeO	H	H	1-Me-6-Ind	V-33,V-34			
V-37	cPenMeO	H	H	2-Ind	V-31,V-32	A	5.35	364(M <sup>+</sup> +1)
V-38	cPenMeO	H	H	1Me-2-Ind	V-29,V-30			
V-39	cPenMeO	H	H	3-Ind	V-31,V-32			
V-40	cPenMeO	H	H	1Me-3-Ind	V-29,V-30	A	4.75	363(M <sup>+</sup> +1)
V-41	cPenMeO	H	H	1iPr-5-Ind	V-29,V-30	C		405(M <sup>+</sup> )
V-42	cPenMeO	H	H	1cPen-5-Ind	V-29,V-30	C		431(M <sup>+</sup> )
V-43	cPenMeO	H	H	1-(2OHEt)-5-Ind	V-43			

Table-V-2

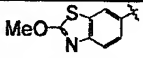

V-44	cPenMeO	Me	H	3CHO-5-Ind	V-44			
V-45	cPenMeO	H	H	3CHO-5-Ind	V-45			
V-46	cPenMeO	H	H	3CHO,1Me-5-Ind	V-29,V-30	C		406(M <sup>+</sup> +1)
V-47	cPenMeO	Me	H	3Ac-5-Ind	V-47			
V-48	cPenMeO	H	H	3Ac-5-Ind	V-48			
V-49	cPenMeO	H	H	3Ac,1Me-5-Ind	V-29,V-30	C		420(M <sup>+</sup> +1)
V-50	cPenMeO	Me	H	3Me-5-Ind	V-50			
V-51	cPenMeO	H	H	3Me-5-Ind	V-51			
V-52	cPenMeO	H	H	1,3DMe-5Ind	V-29,V-30	C		391(M <sup>+</sup> )
V-53	cPenMeO	H	H	1,2,3triMe-5Ind	V-22,V-29,V-30	C		405(M <sup>+</sup> )
V-54	cPenO	Me	H	4-1HIdz	V-54			
V-55	cPenO	H	H	4-1HIdz	V-55			
V-56	cPenO	H	H	1Me-4-1HIdz	V-29,V-30			
V-57	cPenO	Me	H	5-1HIdz	V-57			
V-58	cPenO	H	H	5-1HIdz	V-58			
V-59	cPenO	H	H	1Me-5-1HIdz	V-29,V-30			
V-60	cPenO	H	H	1Et-5-1HIdz	V-29,V-30			
V-61	cPenO	H	H	1Pr-5-1HIdz	V-29,V-30			
V-62	cPenO	H	H	2Me-5-2HIdz	V-29,V-30			
V-63	cPenMeO	H	H	6-1HIdz	V-57,V-58			
V-64	cPenMeO	H	H	1Me-6-1HIdz	V-29,V-30			
V-65	cPenMeO	H	H	1Et-5-1HIdz	V-29,V-30			
V-66	cPenO	Me	H	3Me-5-1HIdz	V-66			
V-67	cPenO	H	H	3Me-5-1HIdz	V-67			
V-68	cPenO	Me	H	1,3DMe-5-1HIdz	V-68			
V-68	cPenO	H	H	1,3DMe-5-1HIdz	V-69			
V-69	cPenO	H	H	3(CHO)-5-1HIdz	V-22,V-23			
V-70	cPenO	H	H	3(OHO),1Me-5-1HIdz	V-22,V-23	A	4.38	365(M <sup>+</sup> +1)
V-71	cPenO	H	H	3OH-5-1HIdz	V-22,V-23			
V-72	cPenO	H	H	3OH,1Me-5-1HIdz	V-22,V-23	A	3.71	381(M <sup>+</sup> +1)
V-73	cPenMeO	Me	H	5-BT	V-73			
V-74	cPenMeO	H	H	5-BT	V-74			
V-75	cPenMeO	H	H	5-BF	V-22,V-23	C		378(M <sup>+</sup> )
V-76	cPenMeO	H	H	2,3DMe-5-BF	V-22,V-23	C		406(M <sup>+</sup> )
V-77	cPenMeO	Me	H	5-2ABzt	V-77			
V-78	cPenMeO	H	H	5-2ABzt	V-78			
V-79	cPenMeO	Et	H	5-Bzt	V-79			
V-80	cPenMeO	H	H	5-Bzt	V-80			
V-81	cPenMeO	Me	H	2Me-5-Bzt	V-81			
V-82	cPenMeO	H	H	2Me-5-Bzt	V-82			
V-83	cPenMeO	Et	H	2,2DMe-5-2ABzt	V-83			
V-84	cPenMeO	H	H	2,2DMe-5-2ABzt	V-84			
V-85	cPenMeO	H	H	6-2ABzt	V-77,V-78	C		397(M <sup>+</sup> +1)
V-86	cPenMeO	H	H	6-Bzt	V-79,V-80	C		453(M <sup>+</sup> +1)
V-87	cPenMeO	H	H	2Me-6-Bzt	V-81,V-82	C		410(M <sup>+</sup> +1)
V-88	cPenMeO	H	H		V-88			
V-89	cPenMeO	H	H		V-89			

Table-V-3

V-90	cPenMeO	H	H		V-29,V-30	C		412(M <sup>+</sup> +1)
V-91	cPenMeO	H	H		V-91	C		414(M <sup>+</sup> +1)
V-92	cPenMeO	H	H		V-29,V-30	C		425(M <sup>+</sup> +1)
V-93	cPenO	H	H		V-22,V-23	B	3.87	368(M <sup>+</sup> +1)
V-94	cPenO	H	H		V-22,V-23	B	3.58	368(M <sup>+</sup> +1)
V-95	cPenO	H	H		V-22,V-23	A	2.57	315(M <sup>+</sup> +1)
V-96	cPenO	H	H		V-22,V-23	A	3.84	351(M <sup>+</sup> +1)
V-97	cPenO	H	H		V-29,V-30	A	4.28	365(M <sup>+</sup> +1)
V-98	cPenMeO	H	H	3-Qu	V-22,V-23	C		376(M <sup>+</sup> +1)
V-99	cPenMeO	H	H	6-Qu	V-22,V-23	C		376(M <sup>+</sup> +1)
V-100	cPenO	H	H	6-IQ	V-22,V-23	A	2.15	452(M <sup>+</sup> +1)
V-101	cPenO	H	H		V-22,V-23	A	3.74	378(M <sup>+</sup> +1)
V-102	cPenMeO	H	H		V-22,V-23	C		378(M <sup>+</sup> +1)
V-103	cHexMeO	Et	H		V-33	C		406(M <sup>+</sup> )
V-104	cHexMeO	H	H		V-34	C		378(M <sup>+</sup> +1)
V-105	cHexMeO	Et	H		V-33	C		422(M <sup>+</sup> )
V-106	cHexMeO	H	H		V-34	C		394(M <sup>+</sup> )
V-107	cHexMeO	H	H		V-22,V-23	C		455(M <sup>+</sup> +1)
V-108	cHexMeO	H	H		V-22,V-23	C		495(M <sup>+</sup> +1)
V-109	cHexMeO	H	H		V-22,V-23	C		487(M <sup>+</sup> +1)
V-110	cPenO	H	H	3(COOH),1Me-7-1HIdz	V-22,V-23	A	3.99	409(M <sup>+</sup> +1)
V-111	cPenO	H	H	3(COOH),1Me-5-1HIdz	V-22,V-23	A	3.75	409(M <sup>+</sup> +1)
V-112	cPenO	H	H	3(COOH),2Me-5-2HIdz	V-22,V-23	A	3.96	409(M <sup>+</sup> +1)
V-113	cPenO	H	H	3(COOH),2Me-7-2HIdz	V-22,V-23	A	3.80	409(M <sup>+</sup> +1)
V-114	cPenO	H	H	3(COOH)-7-1HIdz	V-22,V-23	A	3.66	395(M <sup>+</sup> +1)
V-115	cPenO	H	H	3(COOH)-5-1HIdz	V-22,V-23	A	3.49	395(M <sup>+</sup> +1)

[Examples W-1 to W-25]

Synthesis of 6-bromocinnoline (Intermediate 96)

The title compound (Intermediate 96, 134 mg) was obtained from commercially available 4-bromo-2-iodoaniline (711 mg, Ald) by a method known from literature (Kimball, D. et al., Organic Letter, 2000, p.3825).

#### Synthesis of 7-bromoquinazoline (Intermediate 97)

The title compound (Intermediate 97, 921 mg) was obtained from commercially available quinazoline (2.11 g, WAKO) by a known method described in a publication (Dalby, B. et al., Synthesis, 2002, p.83).

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification are shown in Table-W-1 and Table-W-2.



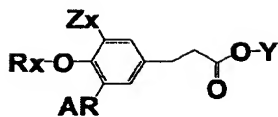
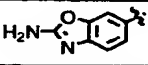
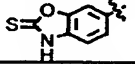
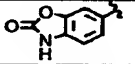

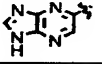
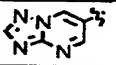
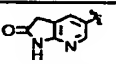


Table-W-1

Exp.	RxO	Y	Zx	AR	Syn	LCMS		
						method	RTime	Mass
W-1	cPenMeO	H	H		V-22,V-23	C		366(M <sup>+</sup> +1)
W-2	cPenMeO	H	H		V-22,V-23	C		383(M <sup>+</sup> +1)
W-3	cPenMeO	H	H		V-22,V-23	C		365(M <sup>+</sup> +1)
W-4	cPenMeO	H	H		V-22,V-23	C		380(M <sup>+</sup> +1)
W-5	cPenMeO	H	H		V-22,V-23	C		366(M <sup>+</sup> +1)
W-6	cPenMeO	H	H		V-22,V-23	C		380(M <sup>+</sup> +1)
W-7	cPenMeO	H	H		V-22,V-23	C		381(M <sup>+</sup> +1)
W-8	cPenMeO	H	H		V-22,V-23	C		398(M <sup>+</sup> +1)
W-9	cPenMeO	H	H		V-22,V-23	C		382(M <sup>+</sup> +1)
W-10	cPenMeO	H	H		V-22,V-23	C		366(M <sup>+</sup> +1)
W-11	cPenMeO	H	H		V-22,V-23	C		377(M <sup>+</sup> +1)
W-12	cPenO	H	H		V-22,V-23	A	3.97	363(M <sup>+</sup> +1)
W-13	cPenO	H	H		V-22,V-23	A	4.06	363(M <sup>+</sup> +1)
W-14	cPenMeO	H	H		V-22,V-23	C		380(M <sup>+</sup> +1)
W-15	cPenO	H	H		V-22,V-23	C		355(M <sup>+</sup> +1)
W-16	cPenMeO	H	H		V-22,V-23	C		397(M <sup>+</sup> +1)
W-17	cPenMeO	H	H		V-22,V-23	C		381(M <sup>+</sup> +1)
W-18	cPenMeO	H	H		V-22,V-23	C		380(M <sup>+</sup> +1)

Table-W-2

W-19	cPenMeO	H	H		V-22,V-23	C		381(M <sup>+</sup> +1)
W-20	cPenMeO	H	H		V-22,V-23	C		398(M <sup>+</sup> +1)
W-21	cPenMeO	H	H		V-22,V-23	C		382(M <sup>+</sup> +1)
W-22	cPenO	H	H		V-22,V-23	C		351(M <sup>+</sup> +1)
W-23	cPenO	H	H		V-22,V-23	C		353(M <sup>+</sup> +1)
W-24	cPenO	H	H		V-22,V-23	C		353(M <sup>+</sup> +1)
W-25	cPenO	H	H		V-22,V-23	C		367(M <sup>+</sup> +1)

## [Example X-1]

Synthesis of ethyl 3-[2-cyclopentyloxy-5-(naphthalen-2-yl)phenyl]acrylate

(Intermediate 98)

According to the procedure described in the synthesis method of Intermediate 7 provided that the reaction was carried out for 1 hour, Compound No. D-20 (396 mg), ethyl diethylphosphonoacetate (288  $\mu$ l), and 60% sodium hydride (59 mg) were reacted and treated to obtain the title compound (Intermediate 98, 428 mg).

Synthesis of ethyl 3-[2-cyclohexylmethyloxy-5-(naphthalen-1-yl)phenyl]propionate (Compound No. X-1)

According to the procedure described in the synthesis method of Intermediate B-99 with the modifications that the reaction was carried out at 50°C for 5 hours, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 10:1), Intermediate 98 (361 mg) and Raney 2800 nickel (380 mg) were reacted and treated to obtain the title compound (Compound No. X-1, 397 mg).

## [Example X-2]

Synthesis of 3-[2-cyclohexylmethoxy-5-(naphthalen-1-yl)phenyl]propionic acid  
(Compound No. X-2)

According to the procedure described in the synthesis method of Intermediate 9 provided that the reaction was carried out for 2.5 hours, Compound No. X-1 (390 mg) and 2 N aqueous sodium hydroxide (1.1 ml) were reacted and treated to obtain the title compound (Compound No. X-2, 338 mg).

[Examples X-1 to X-4]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification including the examples described above are shown in Table-X-1.

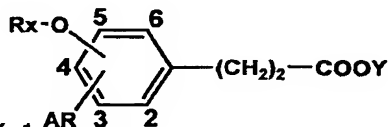


Table-X-1

Exp.	RxO	Y	RxO positio	AR	Syn	AR position	LCMS		
							method	RTime	Mass
X-1	cPenO	Et	2	2-Nap	X-1	5			
X-2	cPenO	H	2	2-Nap	X-2	5	C		347(M <sup>+</sup> +1)
X-3	cPenO	H	2	1Me-5-Ind	X-1,X-2	5	C		350(M <sup>+</sup> +1)
X-4	cPenO	H	2	1Me-5-1HIdz	X-1,X-2	5	C		351(M <sup>+</sup> +1)

[Reference Examples: Intermediates Aa-1 to Aa-47]

Synthesis of methyl 3-[3-(naphthalen-2-yl)-4-trifluoromethanesulfonylphenyl]-propionate (Intermediate Aa-1)

A solution of Intermediate 41 (4.34 g) in dehydrated pyridine (120 ml) was added with trifluoromethanesulfonic anhydride (2.6 ml, ALD) under ice cooling, then warmed to room temperature, and stirred for 4 hours. The reaction mixture was concentrated under reduced pressure, and then extracted with ethyl acetate (800 ml). The organic layer was washed successively with 1 N hydrochloric acid, saturated aqueous ammonium chloride and saturated brine, and dried, and then

the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography (hexane:ethyl acetate = 6:1) to obtain the title compound (Intermediate Aa-1, 4.98 g).

Typical examples of the reaction intermediates including those mentioned above, that can be obtained by reacting and treating corresponding starting compounds according to the synthesis method of Intermediate Aa-1, are shown in Table-Aa-1.

In the column indicated as "Mass" in the table, data of mass spectra measured by fast atom bombardment mass spectrometry (FAB-MS) are shown.

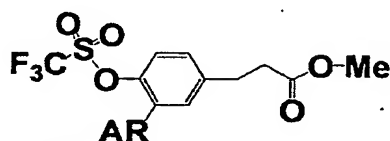


Table-Aa-1

Exp.	AR	Mass	Exp.	AR	Mass
Aa-1	2-Nap	439 ( $M^+ + 1$ )	Aa-25	1Me-4-1HIdz	443 ( $M^+ + 1$ )
Aa-2	5-Ind	428 ( $M^+ + 1$ )	Aa-26	5-1HIdz	429 ( $M^+ + 1$ )
Aa-3	1Me-5-Ind	442 ( $M^+ + 1$ )	Aa-27	1Me-5-1HIdz	443 ( $M^+ + 1$ )
Aa-4	5-1HIdz	429 ( $M^+ + 1$ )	Aa-28	1Et-5-1HIdz	457 ( $M^+ + 1$ )
Aa-5	1Me-5-1HIdz	443 ( $M^+ + 1$ )	Aa-29	1Pr-5-1HIdz	471 ( $M^+ + 1$ )
Aa-6	5-BF	432 ( $M^+ + 1$ )	Aa-30	2Me-5-2HIdz	443 ( $M^+ + 1$ )
Aa-7	3-Qu	440 ( $M^+ + 1$ )	Aa-31	6-1HIdz	429 ( $M^+ + 1$ )
Aa-8	1-Nap	439 ( $M^+ + 1$ )	Aa-32	1Me-6-1HIdz	443 ( $M^+ + 1$ )
Aa-9	6(MeO)-2-Nap	469 ( $M^+ + 1$ )	Aa-33	3Me-5-1HIdz	443 ( $M^+ + 1$ )
Aa-10	6(NMe <sub>2</sub> )-2-Nap	482 ( $M^+ + 1$ )	Aa-34	1,3DMe-5-1HIdz	457 ( $M^+ + 1$ )
Aa-11	4-Ind	428 ( $M^+ + 1$ )	Aa-35	5-BT	445 ( $M^+ + 1$ )
Aa-12	1Me-4-Ind	442 ( $M^+ + 1$ )	Aa-36	2,3DMe-5-BF	457 ( $M^+ + 1$ )
Aa-13	6-Ind	428 ( $M^+ + 1$ )	Aa-37	5-2ABzt	461 ( $M^+ + 1$ )
Aa-14	1Me-6-Ind	442 ( $M^+ + 1$ )	Aa-38	5-Bzt	456 ( $M^+ + 1$ )
Aa-15	2-Ind	428 ( $M^+ + 1$ )	Aa-39	2Me-5-Bzt	460 ( $M^+ + 1$ )
Aa-16	1Me-2-Ind	442 ( $M^+ + 1$ )	Aa-40	2,2DMe-5-2ABzt	489 ( $M^+ + 1$ )
Aa-17	3-Ind	428 ( $M^+ + 1$ )	Aa-41	6-2ABzt	461 ( $M^+ + 1$ )
Aa-18	1Me-3-Ind	442 ( $M^+ + 1$ )	Aa-42	6-Bzt	456 ( $M^+ + 1$ )
Aa-19	1iPr-5-Ind	470 ( $M^+ + 1$ )	Aa-43	2Me-6-Bzt	460 ( $M^+ + 1$ )
Aa-20	1cPen-5-Ind	496 ( $M^+ + 1$ )	Aa-44	6-Qu	440 ( $M^+ + 1$ )
Aa-21	3Me-5-Ind	442 ( $M^+ + 1$ )	Aa-45	6-IQ	440 ( $M^+ + 1$ )
Aa-22	1,3DMe-5Ind	456 ( $M^+ + 1$ )	Aa-46	2-BF	429 ( $M^+ + 1$ )
Aa-23	1,2,3triMe-5Ind	470 ( $M^+ + 1$ )	Aa-47	2-BT	445 ( $M^+ + 1$ )
Aa-24	4-1HIdz	429 ( $M^+ + 1$ )			

[Example Ca-1]

Synthesis of methyl 3-[4-(phenyl)-3-(naphthalen-2-yl)phenyl]propionate (Compound No. Ca-1)

Compound No. Aa-1 (138.4 mg, corresponding to the substance mentioned in the column of SM1 in Table-Ca-1 mentioned later), phenylboronic acid (71.3 mg, corresponding to the substance mentioned in the column of SM 2 mentioned in Table-Ca-1 mentioned later), cesium carbonate (254.9 mg), PdCl<sub>2</sub>(dppf) (25.6 mg) were added with toluene (600  $\mu$ l), methanol (1.2 ml), and water (1.2 ml), and stirred at 80°C for 17 hours under nitrogen atmosphere. The reaction mixture was added with ethyl acetate (30 ml), washed successively with water and saturated brine. The organic layer was dried, and then the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography (hexane:ethyl acetate = 8:1) to obtain the title compound (Compound No. Ca-1, 140.6 mg).

[Example Ca-2]

Synthesis of 3-[4-phenyl-3-(naphthalen-2-yl)phenyl]propionic acid (Compound No. Ca-2)

A solution of Compound Ca-1 (137.7 mg) in methanol (4.0 ml) was added with 2 N aqueous sodium hydroxide (720  $\mu$ l), and stirred at 60°C for 16 hours. The reaction mixture was concentrated under reduced pressure, then made acidic with 5% aqueous hydrochloric acid under ice cooling, and extracted with ethyl acetate (50 ml). The organic layer was washed with saturated brine, and dried, and then the solvent was evaporated under reduced pressure to obtain the title compound (Compound No. Ca-2, 108 mg).

[Examples Ca-1 to Ca-270 and Examples Cb-1 to Cb-95]

Typical examples of the compounds of the present invention including those mentioned in the examples described above, that can be obtained by reacting and treating corresponding starting compounds according to the methods described in

Examples Ca-1 and Ca-2, are shown in Table-Ca-1 to Table-Ca-5, Table-Cb-1 and Table-Cb-2.

The substances mentioned in the columns of "SM1" in the tables correspond to reaction intermediates, and those mentioned in the columns of "SM2" in the tables correspond to the boronic acid reagent used in Example Ca-1. The boronic acid reagents shown with the symbols of "BRA (number)" mentioned in the columns of "SM2" are those mentioned in Table-Ba-1 and Table-Ba-2. The regents for which cells of the columns of "Manufacturer" in the tables are blank are synthesized according to a method described in ordinary chemical literatures.

Table-Ba-1

Reagen	Name of reagent	Manufacturer	Reagen	Name of reagent	Manufacturer
BRA1	Naphthalene-2-boronic acid	TCI	BRA23	Cyclopropyl boronic acid	
BRA2	(1H-Indol-5-yl) boronic acid	Frontier	BRA24	6-Ethoxynaphthalene-2-boronic acid	Ald
BRA3	(1-Methyl-1H-indol-5-yl) boronic acid	Frontier	BRA25	Benzo[b]thiophene-2-boronic acid	LANC
BRA4	(1-Ethyl-1H-indol-5-yl) boronic acid		BRA26	Pyridine-4-boronic acid	ALD
BRA5	(1H-Indazol-5-yl) boronic acid		BRA27	Dibenzofuran-2-boronic acid	Ald
BRA6	(1-Methyl-1H-indazol-5-yl) boronic acid		BRA28	Cyclopentyl boronic acid	LANC
BRA7	(1-Ethyl-1H-indazol-5-yl) boronic acid		BRA29	4-Methylphenyl boronic acid	Ald
BRA8	(2-Methyl-2H-indazol-5-yl) boronic acid		BRA30	4-Chlorophenyl boronic acid	Ald
BRA9	Benzothiazole-6-yl-4,4,5,5-tetramethyl-1,3,2-dioxaborolan		BRA31	1-n-Butyl boronic acid	Ald
BRA10	Quinoline-3-boronic acid	Frontier	BRA32	2-Fluorophenyl boronic acid	Ald
BRA11	Quinoline-6-yl-4,4,5,5-tetramethyl-1,3,2-dioxaborolan	Ald	BRA33	3-Fluorophenyl boronic acid	Ald
BRA12	Isoquinoline-6-yl-4,4,5,5-tetramethyl-1,3,2-dioxaborolan		BRA34	4-Fluorophenyl boronic acid	Ald
BRA13	Methyl boronic acid	Ald	BRA35	2-Furyl boronic acid	Ald
BRA14	Phenyl boronic acid	Ald	BRA36	2-Thienyl boronic acid	Ald
BRA15	4-Hydroxyphenyl boronic acid	Ald	BRA37	3-Methoxyphenyl boronic acid	Ald
BRA16	Naphthalene-1-boronic acid	Ald	BRA38	2-Methoxyphenyl boronic acid	
BRA17	3,5-Bis(trifluoromethyl) phenyl boronic acid	TCI	BRA39	2-(Trifluoromethyl) phenyl boronic acid	
BRA18	Benzo[b]furan-2-boronic acid	Ald	BRA40	3-(Trifluoromethyl) phenyl boronic acid	
BRA19	4-Methoxyphenyl boronic acid	Ald	BRA41	4-(Trifluoromethyl) phenyl boronic acid	
BRA20	2-Methylpropyl boronic acid	Ald	BRA42	Indan-2-yl-4,4,5,5-tetramethyl-1,3,2-dioxaborolane	
BRA21	4-(Dimethylamino) phenyl boronic acid	Ald	BRA43	4-Methylindan-2-yl-4,4,5,5-tetramethyl-1,3,2-dioxaborolane	
BRA22	4-Fluorophenyl boronic acid	TCI	BRA44	5-Methylindan-2-yl-4,4,5,5-tetramethyl-1,3,2-dioxaborolane	

Table-Ba-2

Reagent	Name of reagent	Manufacture	Reagent	Name of reagent	Manufacture
BRA45	4,7-Dimethylindan-2-yl-4,4,5,5-tetramethyl-1,3,2-dioxaborolane		BRA67	3-Furyl boronic acid	Ald
BRA46	5,6-Dimethylindan-2-yl-4,4,5,5-tetramethyl-1,3,2-dioxaborolane		BRA68	3-Thienyl boronic acid	Ald
BRA47	5-Fluoroindan-2-yl-4,4,5,5-tetramethyl-1,3,2-dioxaborolane		BRA69	Pyridine-2-yl-4,4,5,5-tetramethyl-1,3,2-dioxaborolane	
BRA48	4-Fluoroindan-2-yl-4,4,5,5-tetramethyl-1,3,2-dioxaborolane		BRA70	Pyridine-3-boronic acid	Ald
BRA49	4,7-Difluoroindan-2-yl-4,4,5,5-tetramethyl-1,3,2-dioxaborolane		BRA71	2,3-Dimethylphenyl boronic acid	Ald
BRA50	5,6-Difluoroindan-2-yl-4,4,5,5-tetramethyl-1,3,2-dioxaborolane		BRA72	2,5-Dimethylphenyl boronic acid	Ald
BRA51	4-Chloroindan-2-yl-4,4,5,5-tetramethyl-1,3,2-dioxaborolane		BRA73	3,5-Dimethylphenyl boronic acid	Ald
BRA52	5-Chloroindan-2-yl-4,4,5,5-tetramethyl-1,3,2-dioxaborolane		BRA74	2,3-DiChlorophenyl boronic acid	Ald
BRA53	4,7-Dichloroindan-2-yl-4,4,5,5-tetramethyl-1,3,2-dioxaborolane		BRA75	2,4-DiChlorophenyl boronic acid	Ald
BRA54	5,6-Dichloroindan-2-yl-4,4,5,5-tetramethyl-1,3,2-dioxaborolane		BRA76	2,5-DiChlorophenyl boronic acid	Ald
BRA55	4-Methoxyindan-2-yl-4,4,5,5-tetramethyl-1,3,2-dioxaborolane		BRA77	2,6-DiChlorophenyl boronic acid	Acros
BRA56	5-Methoxyindan-2-yl-4,4,5,5-tetramethyl-1,3,2-dioxaborolane		BRA78	3,4-DiChlorophenyl boronic acid	Ald
BRA57	5,6-Dimethoxyindan-2-yl-4,4,5,5-tetramethyl-1,3,2-		BRA79	3,5-DiChlorophenyl boronic acid	Ald
BRA58	Cyclohexyl boronic acid	Ald	BRA80	2,3-Difluorophenyl boronic acid	Ald
BRA59	2-Methylphenyl boronic acid	Ald	BRA81	2,4-Difluorophenyl boronic acid	Ald
BRA60	3-Methylphenyl boronic acid	Ald	BRA82	2,5-Difluorophenyl boronic acid	Ald
BRA61	2-Chlorophenyl boronic acid	Ald	BRA83	2,6-Difluorophenyl boronic acid	Ald
BRA62	3-Chlorophenyl boronic acid	Ald	BRA84	3,4-Difluorophenyl boronic acid	Ald
BRA63	2,3-Bis(trifluoromethyl) phenyl boronic acid		BRA85	3,5-Difluorophenyl boronic acid	Ald
BRA64	2,4-Bis(trifluoromethyl) phenyl boronic acid		BRA86	2-(Dimethylamino) phenyl boronic acid	Digital
BRA65	2,5-Bis(trifluoromethyl) phenyl boronic acid		BRA87	3-(Dimethylamino) phenyl boronic acid	Digital
BRA66	3,4-Bis(trifluoromethyl) phenyl boronic acid		BRA88	4-Phenoxy phenyl boronic acid	Ald



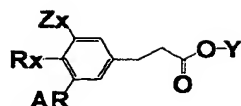


Table-Ca-1

Exp.	Rx	Y	Zx	AR	SM1	SM2	LCMS		
							method	RTime	Mass
Ca-1	Ph	Me	H	2-Nap	Aa-1	BRA14	D		N.D
Ca-2	Ph	H	H	2-Nap	Ca-1	-	C		353 (M <sup>+</sup> +1)
Ca-3	Ph	Me	H	5-Ind	Aa-2	BRA14	C		356 (M <sup>+</sup> +1)
Ca-4	Ph	H	H	5-Ind	Ca-3	-	C		342 (M <sup>+</sup> +1)
Ca-5	Ph	Me	H	1Me-5-Ind	Aa-3	BRA14	C		370 (M <sup>+</sup> +1)
Ca-6	Ph	H	H	1Me-5-Ind	Ca-5	-	C		356 (M <sup>+</sup> +1)
Ca-7	Ph	H	H	5-1HIdz	Aa-4	BRA14	C		343 (M <sup>+</sup> +1)
Ca-8	Ph	Me	H	1Me-5-1HIdz	Aa-5	BRA14	C		371 (M <sup>+</sup> +1)
Ca-9	Ph	H	H	1Me-5-1HIdz	Ca-8	-	C		357 (M <sup>+</sup> +1)
Ca-10	Ph	H	H	5-BF	Aa-6	BRA14	C		342 (M <sup>+</sup> +1)
Ca-11	Ph	H	H	3-Qu	Aa-7	BRA14	C		354 (M <sup>+</sup> +1)
Ca-12	Ph	H	H	1-Nap	Aa-8	BRA14	C		353 (M <sup>+</sup> +1)
Ca-13	Ph	H	H	6(OMe)-2-Nap	Aa-9	BRA14	C		383 (M <sup>+</sup> +1)
Ca-14	Ph	H	H	6(NMe2)-2-Nap	Aa-10	BRA14	C		396 (M <sup>+</sup> +1)
Ca-15	Ph	H	H	4-Ind	Aa-11	BRA14	C		342 (M <sup>+</sup> +1)
Ca-16	Ph	H	H	1Me-4-Ind	Aa-12	BRA14	C		356 (M <sup>+</sup> +1)
Ca-17	Ph	H	H	6-Ind	Aa-13	BRA14	C		342 (M <sup>+</sup> +1)
Ca-18	Ph	H	H	1Me-6-Ind	Aa-14	BRA14	C		356 (M <sup>+</sup> +1)
Ca-19	Ph	H	H	2-Ind	Aa-15	BRA14	C		342 (M <sup>+</sup> +1)
Ca-20	Ph	H	H	1Me-2-Ind	Aa-16	BRA14	C		356 (M <sup>+</sup> +1)
Ca-21	Ph	H	H	3-Ind	Aa-17	BRA14	C		342 (M <sup>+</sup> +1)
Ca-22	Ph	H	H	1Me-3-Ind	Aa-18	BRA14	C		356 (M <sup>+</sup> +1)
Ca-23	Ph	H	H	1iPr-5-Ind	Aa-19	BRA14	C		384 (M <sup>+</sup> +1)
Ca-24	Ph	H	H	1cPen-5-Ind	Aa-20	BRA14	C		410 (M <sup>+</sup> +1)
Ca-25	Ph	H	H	3Me-5-Ind	Aa-21	BRA14	C		356 (M <sup>+</sup> +1)
Ca-26	Ph	H	H	1,3DMe-5Ind	Aa-22	BRA14	C		370 (M <sup>+</sup> +1)
Ca-27	Ph	H	H	1,2,3triMe-5Ind	Aa-23	BRA14	C		384 (M <sup>+</sup> +1)
Ca-28	Ph	H	H	4-1HIdz	Aa-24	BRA14	C		343 (M <sup>+</sup> +1)
Ca-29	Ph	H	H	1Me-4-1HIdz	Aa-25	BRA14	C		357 (M <sup>+</sup> +1)
Ca-30	Ph	H	H	5-1HIdz	Aa-26	BRA14	C		343 (M <sup>+</sup> +1)
Ca-31	Ph	H	H	1Me-5-1HIdz	Aa-27	BRA14	C		357 (M <sup>+</sup> +1)
Ca-32	Ph	H	H	1Et-5-1HIdz	Aa-28	BRA14	C		371 (M <sup>+</sup> +1)
Ca-33	Ph	H	H	1Pr-5-1HIdz	Aa-29	BRA14	C		385 (M <sup>+</sup> +1)
Ca-34	Ph	H	H	2Me-5-2HIdz	Aa-30	BRA14	C		357 (M <sup>+</sup> +1)
Ca-35	Ph	H	H	6-1HIdz	Aa-31	BRA14	C		343 (M <sup>+</sup> +1)
Ca-36	Ph	H	H	1Me-6-1HIdz	Aa-32	BRA14	C		357 (M <sup>+</sup> +1)
Ca-37	Ph	H	H	3Me-5-1HIdz	Aa-33	BRA14	C		357 (M <sup>+</sup> +1)
Ca-38	Ph	H	H	1,3DMe-5-1HIdz	Aa-34	BRA14	C		371 (M <sup>+</sup> +1)
Ca-39	Ph	H	H	5-BT	Aa-35	BRA14	C		359 (M <sup>+</sup> +1)
Ca-40	Ph	H	H	2,3DMe-5-BF	Aa-36	BRA14	C		387 (M <sup>+</sup> +1)
Ca-41	Ph	H	H	5-2ABzt	Aa-37	BRA14	C		375 (M <sup>+</sup> +1)
Ca-42	Ph	H	H	5-Bzt	Aa-38	BRA14	C		360 (M <sup>+</sup> +1)
Ca-43	Ph	H	H	2Me-5-Bzt	Aa-39	BRA14	C		374 (M <sup>+</sup> +1)
Ca-44	Ph	H	H	2,2DMe-5-2ABzt	Aa-40	BRA14	C		403 (M <sup>+</sup> +1)
Ca-45	Ph	H	H	6-2ABzt	Aa-41	BRA14	C		375 (M <sup>+</sup> +1)
Ca-46	Ph	H	H	6-Bzt	Aa-42	BRA14	C		360 (M <sup>+</sup> +1)
Ca-47	Ph	H	H	2Me-6-Bzt	Aa-43	BRA14	C		374 (M <sup>+</sup> +1)
Ca-48	Ph	H	H	6-Qu	Aa-44	BRA14	C		354 (M <sup>+</sup> +1)
Ca-49	Ph	H	H	6-IQ	Aa-45	BRA14	C		354 (M <sup>+</sup> +1)
Ca-50	Ph	H	H	2-BF	Aa-46	BRA14	C		342 (M <sup>+</sup> +1)

Table-Ca-2

Exp.	Rx	Y	Zx	AR	SM1	SM2	LCMS		
							method	RTime	Mass
Ca-51	Ph	H	H	2-BT	Aa-47	BRA14	C		359 (M <sup>+</sup> +1)
Ca-52	4MeOPh	H	H	2-Nap	Aa-1	BRA19	C		383 (M <sup>+</sup> +1)
Ca-53	4MeOPh	H	H	1Me-5-Ind	Aa-3	BRA19	C		386 (M <sup>+</sup> +1)
Ca-54	4MeOPh	H	H	5-1HIdz	Aa-4	BRA19	C		373 (M <sup>+</sup> +1)
Ca-55	4MeOPh	H	H	1Me-5-1HIdz	Aa-5	BRA19	C		387 (M <sup>+</sup> +1)
Ca-56	4MeOPh	H	H	3-Qu	Aa-7	BRA19	C		384 (M <sup>+</sup> +1)
Ca-57	4MeOPh	H	H	1Et-5-1HIdz	Aa-28	BRA19	C		401 (M <sup>+</sup> +1)
Ca-58	3MeOPh	H	H	5-Ind	Aa-2	BRA37	C		372 (M <sup>+</sup> +1)
Ca-59	3MeOPh	H	H	1Me-5-Ind	Aa-3	BRA37	C		386 (M <sup>+</sup> +1)
Ca-60	3MeOPh	H	H	5-1HIdz	Aa-4	BRA37	C		373 (M <sup>+</sup> +1)
Ca-61	3MeOPh	H	H	1Me-5-1HIdz	Aa-5	BRA37	C		387 (M <sup>+</sup> +1)
Ca-62	3MeOPh	H	H	3-Qu	Aa-7	BRA37	C		384 (M <sup>+</sup> +1)
Ca-63	3MeOPh	H	H	1Et-5-1HIdz	Aa-28	BRA37	C		401 (M <sup>+</sup> +1)
Ca-64	2MeOPh	H	H	2-Nap	Aa-1	BRA38	C		383 (M <sup>+</sup> +1)
Ca-65	2MeOPh	H	H	5-Ind	Aa-2	BRA38	C		372 (M <sup>+</sup> +1)
Ca-66	2MeOPh	H	H	1Me-5-Ind	Aa-3	BRA38	C		386 (M <sup>+</sup> +1)
Ca-67	2MeOPh	H	H	5-1HIdz	Aa-4	BRA38	C		373 (M <sup>+</sup> +1)
Ca-68	2MeOPh	H	H	1Me-5-1HIdz	Aa-5	BRA38	C		387 (M <sup>+</sup> +1)
Ca-69	2MeOPh	H	H	5-Bzt	Aa-38	BRA38	C		390 (M <sup>+</sup> +1)
Ca-70	2MeOPh	H	H	3-Qu	Aa-7	BRA38	C		384 (M <sup>+</sup> +1)
Ca-71	2MeOPh	H	H	1Et-5-1HIdz	Aa-28	BRA38	C		401 (M <sup>+</sup> +1)
Ca-72	2MePh	H	H	2-Nap	Aa-1	BRA59	C		367 (M <sup>+</sup> +1)
Ca-73	2MePh	H	H	5-Ind	Aa-2	BRA59	C		356 (M <sup>+</sup> +1)
Ca-74	2MePh	H	H	1Me-5-Ind	Aa-3	BRA59	C		370 (M <sup>+</sup> +1)
Ca-75	2MePh	H	H	5-1HIdz	Aa-4	BRA59	C		357 (M <sup>+</sup> +1)
Ca-76	2MePh	H	H	1Me-5-1HIdz	Aa-5	BRA59	C		371 (M <sup>+</sup> +1)
Ca-77	2MePh	H	H	5-Bzt	Aa-38	BRA59	C		374 (M <sup>+</sup> +1)
Ca-78	3MePh	H	H	2-Nap	Aa-1	BRA60	C		367 (M <sup>+</sup> +1)
Ca-79	3MePh	H	H	5-Ind	Aa-2	BRA60	C		356 (M <sup>+</sup> +1)
Ca-80	3MePh	H	H	5-1HIdz	Aa-4	BRA60	C		357 (M <sup>+</sup> +1)
Ca-81	3MePh	H	H	1Me-5-1HIdz	Aa-5	BRA60	C		371 (M <sup>+</sup> +1)
Ca-82	3MePh	H	H	5-Bzt	Aa-38	BRA60	C		374 (M <sup>+</sup> +1)
Ca-83	3MePh	H	H	1Et-5-1HIdz	Aa-28	BRA60	C		385 (M <sup>+</sup> +1)
Ca-84	4MePh	H	H	2-Nap	Aa-1	BRA29	C		367 (M <sup>+</sup> +1)
Ca-85	4MePh	H	H	5-Ind	Aa-2	BRA29	C		356 (M <sup>+</sup> +1)
Ca-86	4MePh	H	H	1Me-5-Ind	Aa-3	BRA29	C		370 (M <sup>+</sup> +1)
Ca-87	4MePh	H	H	5-1HIdz	Aa-4	BRA29	C		357 (M <sup>+</sup> +1)
Ca-88	4MePh	H	H	1Me-5-1HIdz	Aa-5	BRA29	C		371 (M <sup>+</sup> +1)
Ca-89	4MePh	H	H	5-Bzt	Aa-38	BRA29	C		374 (M <sup>+</sup> +1)
Ca-90	4MePh	H	H	3-Qu	Aa-7	BRA29	C		368 (M <sup>+</sup> +1)
Ca-91	4MePh	H	H	1Et-5-1HIdz	Aa-28	BRA29	C		385 (M <sup>+</sup> +1)
Ca-92	2,3DMePh	H	H	5-Ind	Aa-2	BRA71	C		370 (M <sup>+</sup> +1)
Ca-93	2,3DMePh	H	H	1Me-5-Ind	Aa-3	BRA71	C		384 (M <sup>+</sup> +1)
Ca-94	2,3DMePh	H	H	5-1HIdz	Aa-4	BRA71	C		371 (M <sup>+</sup> +1)
Ca-95	2,3DMePh	H	H	1Me-5-1HIdz	Aa-5	BRA71	C		385 (M <sup>+</sup> +1)
Ca-96	2,3DMePh	H	H	1Et-5-1HIdz	Aa-28	BRA71	C		399 (M <sup>+</sup> +1)
Ca-97	2,5DMePh	H	H	2-Nap	Aa-1	BRA72	C		381 (M <sup>+</sup> +1)
Ca-98	2,5DMePh	H	H	1Me-5-Ind	Aa-3	BRA72	C		384 (M <sup>+</sup> +1)
Ca-99	2,5DMePh	H	H	5-1HIdz	Aa-4	BRA72	C		371 (M <sup>+</sup> +1)
Ca-100	2,5DMePh	H	H	1Me-5-1HIdz	Aa-5	BRA72	C		385 (M <sup>+</sup> +1)
Ca-101	2,5DMePh	H	H	1Et-5-1HIdz	Aa-28	BRA72	C		399 (M <sup>+</sup> +1)
Ca-102	3,5DMePh	H	H	2-Nap	Aa-1	BRA73	C		381 (M <sup>+</sup> +1)
Ca-103	3,5DMePh	H	H	1Me-5-Ind	Aa-3	BRA73	C		384 (M <sup>+</sup> +1)
Ca-104	3,5DMePh	H	H	1Me-5-1HIdz	Aa-5	BRA73	C		385 (M <sup>+</sup> +1)
Ca-105	2CF <sub>3</sub> Ph	H	H	2-Nap	Aa-1	BRA39	C		421 (M <sup>+</sup> +1)

Table-Ca-3

Exp.	Rx	Y	Zx	AR	SM1	SM2	LCMS		
							method	RTime	Mass
Ca-106	2CF <sub>3</sub> Ph	H	H	5-Ind	Aa-2	BRA39	C		410 (M <sup>+</sup> +1)
Ca-107	2CF <sub>3</sub> Ph	H	H	1Me-5-1HIdz	Aa-5	BRA39	C		425 (M <sup>+</sup> +1)
Ca-108	2CF <sub>3</sub> Ph	H	H	5-Bzt	Aa-38	BRA39	C		428 (M <sup>+</sup> +1)
Ca-109	2CF <sub>3</sub> Ph	H	H	3-Qu	Aa-7	BRA39	C		422 (M <sup>+</sup> +1)
Ca-110	2CF <sub>3</sub> Ph	H	H	1Et-5-1HIdz	Aa-28	BRA39	C		439 (M <sup>+</sup> +1)
Ca-111	3CF <sub>3</sub> Ph	H	H	2-Nap	Aa-1	BRA40	C		421 (M <sup>+</sup> +1)
Ca-112	3CF <sub>3</sub> Ph	H	H	5-Ind	Aa-2	BRA40	C		410 (M <sup>+</sup> +1)
Ca-113	3CF <sub>3</sub> Ph	H	H	1Me-5-Ind	Aa-3	BRA40	C		424 (M <sup>+</sup> +1)
Ca-114	3CF <sub>3</sub> Ph	H	H	1Me-5-1HIdz	Aa-5	BRA40	C		425 (M <sup>+</sup> +1)
Ca-115	3CF <sub>3</sub> Ph	H	H	5-Bzt	Aa-38	BRA40	C		428 (M <sup>+</sup> +1)
Ca-116	3CF <sub>3</sub> Ph	H	H	3-Qu	Aa-7	BRA40	C		422 (M <sup>+</sup> +1)
Ca-117	4CF <sub>3</sub> Ph	H	H	5-Ind	Aa-2	BRA41	C		410 (M <sup>+</sup> +1)
Ca-118	4CF <sub>3</sub> Ph	H	H	5-1HIdz	Aa-4	BRA41	C		411 (M <sup>+</sup> +1)
Ca-119	4CF <sub>3</sub> Ph	H	H	1Me-5-1HIdz	Aa-5	BRA41	C		425 (M <sup>+</sup> +1)
Ca-120	4CF <sub>3</sub> Ph	H	H	5-Bzt	Aa-38	BRA41	C		428 (M <sup>+</sup> +1)
Ca-121	4CF <sub>3</sub> Ph	H	H	3-Qu	Aa-7	BRA41	C		422 (M <sup>+</sup> +1)
Ca-122	4CF <sub>3</sub> Ph	H	H	1Et-5-1HIdz	Aa-28	BRA41	C		439 (M <sup>+</sup> +1)
Ca-123	2ClPh	H	H	5-Ind	Aa-2	BRA61	C		376 (M <sup>+</sup> +1)
Ca-124	2ClPh	H	H	5-1HIdz	Aa-4	BRA61	C		377 (M <sup>+</sup> +1)
Ca-125	2ClPh	H	H	1Me-5-1HIdz	Aa-5	BRA61	C		391 (M <sup>+</sup> +1)
Ca-126	2ClPh	H	H	3-Qu	Aa-7	BRA61	C		388 (M <sup>+</sup> +1)
Ca-127	3ClPh	H	H	2-Nap	Aa-1	BRA62	C		387 (M <sup>+</sup> +1)
Ca-128	3ClPh	H	H	1Me-5-Ind	Aa-3	BRA62	C		390 (M <sup>+</sup> +1)
Ca-129	3ClPh	H	H	5-1HIdz	Aa-4	BRA62	C		377 (M <sup>+</sup> +1)
Ca-130	3ClPh	H	H	1Me-5-1HIdz	Aa-5	BRA62	C		391 (M <sup>+</sup> +1)
Ca-131	3ClPh	H	H	5-Bzt	Aa-38	BRA62	C		394 (M <sup>+</sup> +1)
Ca-132	4ClPh	H	H	5-Ind	Aa-2	BRA30	C		376 (M <sup>+</sup> +1)
Ca-133	4ClPh	H	H	1Me-5-Ind	Aa-3	BRA30	C		390 (M <sup>+</sup> +1)
Ca-134	4ClPh	H	H	1Me-5-1HIdz	Aa-5	BRA30	C		391 (M <sup>+</sup> +1)
Ca-135	4ClPh	H	H	5-Bzt	Aa-38	BRA30	C		394 (M <sup>+</sup> +1)
Ca-136	2,3DCIPh	H	H	5-Ind	Aa-2	BRA74	C		411 (M <sup>+</sup> +1)
Ca-137	2,3DCIPh	H	H	1Me-5-Ind	Aa-3	BRA74	C		425 (M <sup>+</sup> +1)
Ca-138	2,3DCIPh	H	H	1Me-5-1HIdz	Aa-5	BRA74	C		426 (M <sup>+</sup> +1)
Ca-139	2,4DCIPh	H	H	5-Ind	Aa-2	BRA75	C		411 (M <sup>+</sup> +1)
Ca-140	2,4DCIPh	H	H	1Me-5-1HIdz	Aa-5	BRA75	C		426 (M <sup>+</sup> +1)
Ca-141	2,4DCIPh	H	H	5-Bzt	Aa-38	BRA75	C		429 (M <sup>+</sup> +1)
Ca-142	2,5DCIPh	H	H	1Me-5-Ind	Aa-3	BRA76	C		425 (M <sup>+</sup> +1)
Ca-143	2,5DCIPh	H	H	1Me-5-1HIdz	Aa-5	BRA76	C		426 (M <sup>+</sup> +1)
Ca-144	2,6DCIPh	H	H	1Me-5-1HIdz	Aa-5	BRA77	C		426 (M <sup>+</sup> +1)
Ca-145	3,4DCIPh	H	H	2-Nap	Aa-1	BRA78	C		421 (M <sup>+</sup> +1)
Ca-146	3,4DCIPh	H	H	5-Ind	Aa-2	BRA78	C		411 (M <sup>+</sup> +1)
Ca-147	3,4DCIPh	H	H	1Me-5-Ind	Aa-3	BRA78	C		425 (M <sup>+</sup> +1)
Ca-148	3,4DCIPh	H	H	1Me-5-1HIdz	Aa-5	BRA78	C		426 (M <sup>+</sup> +1)
Ca-149	3,5DCIPh	H	H	2-Nap	Aa-1	BRA79	C		421 (M <sup>+</sup> +1)
Ca-150	3,5DCIPh	H	H	1Me-5-Ind	Aa-3	BRA79	C		425 (M <sup>+</sup> +1)
Ca-151	3,5DCIPh	H	H	1Me-5-1HIdz	Aa-5	BRA79	C		426 (M <sup>+</sup> +1)
Ca-152	2FPh	H	H	2-Nap	Aa-1	BRA32	C		371 (M <sup>+</sup> +1)
Ca-153	2FPh	H	H	1Me-5-Ind	Aa-3	BRA32	C		374 (M <sup>+</sup> +1)
Ca-154	2FPh	H	H	5-1HIdz	Aa-4	BRA32	C		361 (M <sup>+</sup> +1)
Ca-155	2FPh	H	H	1Me-5-1HIdz	Aa-5	BRA32	C		375 (M <sup>+</sup> +1)
Ca-156	2FPh	H	H	5-Bzt	Aa-38	BRA32	C		378 (M <sup>+</sup> +1)
Ca-157	2FPh	H	H	3-Qu	Aa-7	BRA32	C		372 (M <sup>+</sup> +1)
Ca-158	3FPh	H	H	5-Ind	Aa-2	BRA33	C		360 (M <sup>+</sup> +1)
Ca-159	3FPh	H	H	5-1HIdz	Aa-4	BRA33	C		361 (M <sup>+</sup> +1)
Ca-160	3FPh	H	H	1Me-5-1HIdz	Aa-5	BRA33	C		375 (M <sup>+</sup> +1)

Table-Ca-4

Exp.	Rx	Y	Zx	AR	SM1	SM2	LCMS		
							method	RTime	Mass
Ca-161	3FPh	H	H	3-Qu	Aa-7	BRA33	C		372 (M <sup>+</sup> +1)
Ca-162	4FPh	H	H	2-Nap	Aa-1	BRA34	C		371 (M <sup>+</sup> +1)
Ca-163	4FPh	H	H	5-Ind	Aa-2	BRA34	C		360 (M <sup>+</sup> +1)
Ca-164	4FPh	H	H	5-1HIdz	Aa-4	BRA34	C		361 (M <sup>+</sup> +1)
Ca-165	4FPh	H	H	1Me-5-1HIdz	Aa-5	BRA34	C		375 (M <sup>+</sup> +1)
Ca-166	4FPh	H	H	3-Qu	Aa-7	BRA34	C		372 (M <sup>+</sup> +1)
Ca-167	2,3DFPh	H	H	2-Nap	Aa-1	BRA80	C		389 (M <sup>+</sup> +1)
Ca-168	2,3DFPh	H	H	5-Ind	Aa-2	BRA80	C		378 (M <sup>+</sup> +1)
Ca-169	2,3DFPh	H	H	1Me-5-1HIdz	Aa-5	BRA80	C		393 (M <sup>+</sup> +1)
Ca-170	2,4DFPh	H	H	2-Nap	Aa-1	BRA81	C		389 (M <sup>+</sup> +1)
Ca-171	2,4DFPh	H	H	5-Ind	Aa-2	BRA81	C		378 (M <sup>+</sup> +1)
Ca-172	2,4DFPh	H	H	1Me-5-Ind	Aa-3	BRA81	C		392 (M <sup>+</sup> +1)
Ca-173	2,4DFPh	H	H	1Me-5-1HIdz	Aa-5	BRA81	C		393 (M <sup>+</sup> +1)
Ca-174	2,5DFPh	H	H	2-Nap	Aa-1	BRA82	C		389 (M <sup>+</sup> +1)
Ca-175	2,5DFPh	H	H	1Me-5-Ind	Aa-3	BRA82	C		392 (M <sup>+</sup> +1)
Ca-176	2,5DFPh	H	H	1Me-5-1HIdz	Aa-5	BRA82	C		393 (M <sup>+</sup> +1)
Ca-177	2,6DFPh	H	H	2-Nap	Aa-1	BRA83	C		389 (M <sup>+</sup> +1)
Ca-178	2,6DFPh	H	H	1Me-5-Ind	Aa-3	BRA83	C		392 (M <sup>+</sup> +1)
Ca-179	2,6DFPh	H	H	5-1HIdz	Aa-4	BRA83	C		379 (M <sup>+</sup> +1)
Ca-180	2,6DFPh	H	H	1Me-5-1HIdz	Aa-5	BRA83	C		393 (M <sup>+</sup> +1)
Ca-181	3,4DFPh	H	H	2-Nap	Aa-1	BRA84	C		389 (M <sup>+</sup> +1)
Ca-182	3,4DFPh	H	H	5-Ind	Aa-2	BRA84	C		378 (M <sup>+</sup> +1)
Ca-183	3,4DFPh	H	H	1Me-5-1HIdz	Aa-5	BRA84	C		393 (M <sup>+</sup> +1)
Ca-184	3,5DFPh	H	H	2-Nap	Aa-1	BRA85	C		389 (M <sup>+</sup> +1)
Ca-185	3,5DFPh	H	H	1Me-5-Ind	Aa-3	BRA85	C		392 (M <sup>+</sup> +1)
Ca-186	3,5DFPh	H	H	5-1HIdz	Aa-4	BRA85	C		379 (M <sup>+</sup> +1)
Ca-187	3,5DFPh	H	H	1Me-5-1HIdz	Aa-5	BRA85	C		393 (M <sup>+</sup> +1)
Ca-188	2,3(CF <sub>3</sub> ) <sub>2</sub> Ph	H	H	2-Nap	Aa-1	BRA63	C		489 (M <sup>+</sup> +1)
Ca-189	2,3(CF <sub>3</sub> ) <sub>2</sub> Ph	H	H	1Me-5-Ind	Aa-3	BRA63	C		492 (M <sup>+</sup> +1)
Ca-190	2,3(CF <sub>3</sub> ) <sub>2</sub> Ph	H	H	1Me-5-1HIdz	Aa-5	BRA63	C		493 (M <sup>+</sup> +1)
Ca-191	2,4(CF <sub>3</sub> ) <sub>2</sub> Ph	H	H	2-Nap	Aa-1	BRA64	C		489 (M <sup>+</sup> +1)
Ca-192	2,4(CF <sub>3</sub> ) <sub>2</sub> Ph	H	H	1Me-5-1HIdz	Aa-5	BRA64	C		493 (M <sup>+</sup> +1)
Ca-193	2,5(CF <sub>3</sub> ) <sub>2</sub> Ph	H	H	2-Nap	Aa-1	BRA65	C		489 (M <sup>+</sup> +1)
Ca-194	2,5(CF <sub>3</sub> ) <sub>2</sub> Ph	H	H	5-Ind	Aa-2	BRA65	C		478 (M <sup>+</sup> +1)
Ca-195	2,5(CF <sub>3</sub> ) <sub>2</sub> Ph	H	H	1Me-5-1HIdz	Aa-5	BRA65	C		493 (M <sup>+</sup> +1)
Ca-196	2,5(CF <sub>3</sub> ) <sub>2</sub> Ph	H	H	3-Qu	Aa-7	BRA65	C		490 (M <sup>+</sup> +1)
Ca-197	3,4(CF <sub>3</sub> ) <sub>2</sub> Ph	H	H	2-Nap	Aa-1	BRA66	C		489 (M <sup>+</sup> +1)
Ca-198	3,4(CF <sub>3</sub> ) <sub>2</sub> Ph	H	H	1Me-5-Ind	Aa-3	BRA66	C		492 (M <sup>+</sup> +1)
Ca-199	3,4(CF <sub>3</sub> ) <sub>2</sub> Ph	H	H	5-1HIdz	Aa-4	BRA66	C		479 (M <sup>+</sup> +1)
Ca-200	3,4(CF <sub>3</sub> ) <sub>2</sub> Ph	H	H	1Me-5-1HIdz	Aa-5	BRA66	C		493 (M <sup>+</sup> +1)
Ca-201	3,5(CF <sub>3</sub> ) <sub>2</sub> Ph	H	H	5-Ind	Aa-2	BRA17	C		478 (M <sup>+</sup> +1)
Ca-202	3,5(CF <sub>3</sub> ) <sub>2</sub> Ph	H	H	5-1HIdz	Aa-4	BRA17	C		479 (M <sup>+</sup> +1)
Ca-203	3,5(CF <sub>3</sub> ) <sub>2</sub> Ph	H	H	1Me-5-1HIdz	Aa-5	BRA17	C		493 (M <sup>+</sup> +1)
Ca-204	2-Furyl	H	H	2-Nap	Aa-1	BRA35	C		343 (M <sup>+</sup> +1)
Ca-205	2-Furyl	H	H	5-Ind	Aa-2	BRA35	C		332 (M <sup>+</sup> +1)
Ca-206	2-Furyl	H	H	1Me-5-1HIdz	Aa-5	BRA35	C		347 (M <sup>+</sup> +1)
Ca-207	2-Furyl	H	H	3-Qu	Aa-7	BRA35	C		344 (M <sup>+</sup> +1)
Ca-208	3-Furyl	H	H	1Me-5-Ind	Aa-3	BRA67	C		346 (M <sup>+</sup> +1)
Ca-209	3-Furyl	H	H	5-1HIdz	Aa-4	BRA67	C		333 (M <sup>+</sup> +1)
Ca-210	3-Furyl	H	H	1Me-5-1HIdz	Aa-5	BRA67	C		347 (M <sup>+</sup> +1)
Ca-211	2-Thienyl	H	H	2-Nap	Aa-1	BRA36	C		359 (M <sup>+</sup> +1)
Ca-212	2-Thienyl	H	H	1Me-5-Ind	Aa-3	BRA36	C		362 (M <sup>+</sup> +1)
Ca-213	2-Thienyl	H	H	1Me-5-1HIdz	Aa-5	BRA36	C		363 (M <sup>+</sup> +1)
Ca-214	2-Thienyl	H	H	1Et-5-1HIdz	Aa-28	BRA36	C		377 (M <sup>+</sup> +1)
Ca-215	3-Thienyl	H	H	5-Ind	Aa-2	BRA68	C		348 (M <sup>+</sup> +1)

Table-Ca-5

Exp.	Rx	Y	Zx	AR	SM1	SM2	LCMS		
							method	RTime	Mass
Ca-216	3-Thienyl	H	H	1Me-5-Ind	Aa-3	BRA68	C		362 (M <sup>+</sup> +1)
Ca-217	3-Thienyl	H	H	5-1HIdz	Aa-4	BRA68	C		349 (M <sup>+</sup> +1)
Ca-218	3-Thienyl	H	H	1Me-5-1HIdz	Aa-5	BRA68	C		363 (M <sup>+</sup> +1)
Ca-219	3-Thienyl	H	H	5-Bzt	Aa-38	BRA68	C		366 (M <sup>+</sup> +1)
Ca-220	3-Thienyl	H	H	3-Qu	Aa-7	BRA68	C		360 (M <sup>+</sup> +1)
Ca-221	3-Thienyl	H	H	1Et-5-1HIdz	Aa-28	BRA68	C		377 (M <sup>+</sup> +1)
Ca-222	2-Py	H	H	5-Ind	Aa-2	BRA69	C		343 (M <sup>+</sup> +1)
Ca-223	2-Py	H	H	1Me-5-1HIdz	Aa-5	BRA69	C		358 (M <sup>+</sup> +1)
Ca-224	2-Py	H	H	5-Bzt	Aa-38	BRA69	C		361 (M <sup>+</sup> +1)
Ca-225	3-Py	H	H	2-Nap	Aa-1	BRA70	C		354 (M <sup>+</sup> +1)
Ca-226	3-Py	H	H	5-Ind	Aa-2	BRA70	C		343 (M <sup>+</sup> +1)
Ca-227	3-Py	H	H	1Me-5-Ind	Aa-3	BRA70	C		357 (M <sup>+</sup> +1)
Ca-228	3-Py	H	H	1Me-5-1HIdz	Aa-5	BRA70	C		358 (M <sup>+</sup> +1)
Ca-229	3-Py	H	H	1Et-5-1HIdz	Aa-28	BRA70	C		372 (M <sup>+</sup> +1)
Ca-230	4-Py	H	H	2-Nap	Aa-1	BRA26	C		354 (M <sup>+</sup> +1)
Ca-231	4-Py	H	H	5-Ind	Aa-2	BRA26	C		343 (M <sup>+</sup> +1)
Ca-232	4-Py	H	H	1Me-5-Ind	Aa-3	BRA26	C		357 (M <sup>+</sup> +1)
Ca-233	4-Py	H	H	5-1HIdz	Aa-4	BRA26	C		344 (M <sup>+</sup> +1)
Ca-234	4-Py	H	H	1Me-5-1HIdz	Aa-5	BRA26	C		358 (M <sup>+</sup> +1)
Ca-235	4-Py	H	H	5-Bzt	Aa-38	BRA26	C		361 (M <sup>+</sup> +1)
Ca-236	4-Py	H	H	3-Qu	Aa-7	BRA26	C		355 (M <sup>+</sup> +1)
Ca-237	4-Py	H	H	1Et-5-1HIdz	Aa-28	BRA26	C		372 (M <sup>+</sup> +1)
Ca-238	2DMAPh	H	H	2-Nap	Aa-1	BRA86	C		396 (M <sup>+</sup> +1)
Ca-239	2DMAPh	H	H	5-Ind	Aa-2	BRA86	C		385 (M <sup>+</sup> +1)
Ca-240	2DMAPh	H	H	1Me-5-1HIdz	Aa-5	BRA86	C		400 (M <sup>+</sup> +1)
Ca-241	2DMAPh	H	H	5-Bzt	Aa-38	BRA86	C		403 (M <sup>+</sup> +1)
Ca-242	2DMAPh	H	H	1Et-5-1HIdz	Aa-28	BRA86	C		414 (M <sup>+</sup> +1)
Ca-243	3DMAPh	H	H	1Me-5-Ind	Aa-3	BRA87	C		399 (M <sup>+</sup> +1)
Ca-244	3DMAPh	H	H	5-1HIdz	Aa-4	BRA87	C		386 (M <sup>+</sup> +1)
Ca-245	3DMAPh	H	H	1Me-5-1HIdz	Aa-5	BRA87	C		400 (M <sup>+</sup> +1)
Ca-246	3DMAPh	H	H	5-Bzt	Aa-38	BRA87	C		403 (M <sup>+</sup> +1)
Ca-247	3DMAPh	H	H	3-Qu	Aa-7	BRA87	C		397 (M <sup>+</sup> +1)
Ca-248	4DMAPh	H	H	2-Nap	Aa-1	BRA21	C		396 (M <sup>+</sup> +1)
Ca-249	4DMAPh	H	H	1Me-5-Ind	Aa-3	BRA21	C		399 (M <sup>+</sup> +1)
Ca-250	4DMAPh	H	H	1Me-5-1HIdz	Aa-5	BRA21	C		400 (M <sup>+</sup> +1)
Ca-251	4DMAPh	H	H	3-Qu	Aa-7	BRA21	C		397 (M <sup>+</sup> +1)
Ca-252	4DMAPh	H	H	1Et-5-1HIdz	Aa-28	BRA21	C		414 (M <sup>+</sup> +1)
Ca-253	1-Nap	H	H	2-Nap	Aa-1	BRA16	C		403 (M <sup>+</sup> +1)
Ca-254	1-Nap	H	H	5-Ind	Aa-2	BRA16	C		392 (M <sup>+</sup> +1)
Ca-255	1-Nap	H	H	1Me-5-1HIdz	Aa-5	BRA16	C		407 (M <sup>+</sup> +1)
Ca-256	1-Nap	H	H	5-Bzt	Aa-38	BRA16	C		410 (M <sup>+</sup> +1)
Ca-257	2-Nap	H	H	2-Nap	Aa-1	BRA1	C		403 (M <sup>+</sup> +1)
Ca-258	2-Nap	H	H	1Me-5-Ind	Aa-3	BRA1	C		406 (M <sup>+</sup> +1)
Ca-259	2-Nap	H	H	5-1HIdz	Aa-4	BRA1	C		393 (M <sup>+</sup> +1)
Ca-260	2-Nap	H	H	1Me-5-1HIdz	Aa-5	BRA1	C		407 (M <sup>+</sup> +1)
Ca-261	2-Nap	H	H	5-Bzt	Aa-38	BRA1	C		410 (M <sup>+</sup> +1)
Ca-262	5-Ind	H	H	2-Nap	Aa-1	BRA2	C		392 (M <sup>+</sup> +1)
Ca-263	5-Ind	H	H	5-Ind	Aa-2	BRA2	C		381 (M <sup>+</sup> +1)
Ca-264	5-Ind	H	H	1Me-5-1HIdz	Aa-5	BRA2	C		396 (M <sup>+</sup> +1)
Ca-265	5-Ind	H	H	3-Qu	Aa-7	BRA2	C		393 (M <sup>+</sup> +1)
Ca-266	5-Ind	H	H	1Et-5-1HIdz	Aa-28	BRA2	C		410 (M <sup>+</sup> +1)
Ca-267	1Me-5-1HIdz	H	H	2-Nap	Aa-1	BRA6	C		407 (M <sup>+</sup> +1)
Ca-268	1Me-5-1HIdz	H	H	1Me-5-Ind	Aa-3	BRA6	C		410 (M <sup>+</sup> +1)
Ca-269	1Me-5-1HIdz	H	H	1Me-5-1HIdz	Aa-5	BRA6	C		411 (M <sup>+</sup> +1)
Ca-270	1Me-5-1HIdz	H	H	5-Bzt	Aa-38	BRA6	C		414 (M <sup>+</sup> +1)

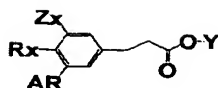


Table-Cb-1

Exp.	Rx	Y	Zx	AR	SM1	SM2	LCMS		
							method	RTime	Mass
Cb-1	cPen	H	H	2-Nap	Aa-1	BRA28	C		345 (M <sup>+</sup> +1)
Cb-2	cPen	H	H	5-Ind	Aa-2	BRA28	C		334 (M <sup>+</sup> +1)
Cb-3	cPen	H	H	1Me-5-Ind	Aa-3	BRA28	C		348 (M <sup>+</sup> +1)
Cb-4	cPen	H	H	5-1HIdz	Aa-4	BRA28	C		335 (M <sup>+</sup> +1)
Cb-5	cPen	H	H	1Me-5-1HIdz	Aa-5	BRA28	C		349 (M <sup>+</sup> +1)
Cb-6	cPen	H	H	5-Bzt	Aa-38	BRA28	C		352 (M <sup>+</sup> +1)
Cb-7	cPen	H	H	3-Qu	Aa-7	BRA28	C		346 (M <sup>+</sup> +1)
Cb-8	cPen	H	H	1Et-5-1HIdz	Aa-28	BRA28	C		363 (M <sup>+</sup> +1)
Cb-9	nBu	H	H	2-Nap	Aa-1	BRA31	C		333 (M <sup>+</sup> +1)
Cb-10	nBu	H	H	5-Ind	Aa-2	BRA31	C		322 (M <sup>+</sup> +1)
Cb-11	nBu	H	H	1Me-5-1HIdz	Aa-5	BRA31	C		337 (M <sup>+</sup> +1)
Cb-12	iBu	H	H	2-Nap	Aa-1	BRA20	C		333 (M <sup>+</sup> +1)
Cb-13	iBu	H	H	1Me-5-Ind	Aa-3	BRA20	C		336 (M <sup>+</sup> +1)
Cb-14	iBu	H	H	1Me-5-1HIdz	Aa-5	BRA20	C		337 (M <sup>+</sup> +1)
Cb-15	iBu	H	H	5-Bzt	Aa-38	BRA20	C		340 (M <sup>+</sup> +1)
Cb-16	iBu	H	H	1Et-5-1HIdz	Aa-28	BRA20	C		351 (M <sup>+</sup> +1)
Cb-17	2-Indan	H	H	2-Nap	Aa-1	BRA42	C		393 (M <sup>+</sup> +1)
Cb-18	2-Indan	H	H	5-Ind	Aa-2	BRA42	C		382 (M <sup>+</sup> +1)
Cb-19	2-Indan	H	H	1Me-5-Ind	Aa-3	BRA42	C		396 (M <sup>+</sup> +1)
Cb-20	2-Indan	H	H	5-1HIdz	Aa-4	BRA42	C		382 (M <sup>+</sup> +1)
Cb-21	2-Indan	H	H	1Me-5-1HIdz	Aa-5	BRA42	C		397 (M <sup>+</sup> +1)
Cb-22	2-Indan	H	H	5-Bzt	Aa-38	BRA42	C		400 (M <sup>+</sup> +1)
Cb-23	2-Indan	H	H	3-Qu	Aa-7	BRA42	C		394 (M <sup>+</sup> +1)
Cb-24	2-Indan	H	H	1Et-5-1HIdz	Aa-28	BRA42	C		411 (M <sup>+</sup> +1)
Cb-25	4Me-2-Indan	H	H	5-Ind	Aa-2	BRA43	C		396 (M <sup>+</sup> +1)
Cb-26	4Me-2-Indan	H	H	5-1HIdz	Aa-4	BRA43	C		397 (M <sup>+</sup> +1)
Cb-27	4Me-2-Indan	H	H	1Me-5-1HIdz	Aa-5	BRA43	C		411 (M <sup>+</sup> +1)
Cb-28	4Me-2-Indan	H	H	3-Qu	Aa-7	BRA43	C		408 (M <sup>+</sup> +1)
Cb-29	5Me-2-Indan	H	H	2-Nap	Aa-1	BRA44	C		407 (M <sup>+</sup> +1)
Cb-30	5Me-2-Indan	H	H	5-Ind	Aa-2	BRA44	C		396 (M <sup>+</sup> +1)
Cb-31	5Me-2-Indan	H	H	5-1HIdz	Aa-4	BRA44	C		397 (M <sup>+</sup> +1)
Cb-32	5Me-2-Indan	H	H	1Me-5-1HIdz	Aa-5	BRA44	C		411 (M <sup>+</sup> +1)
Cb-33	5Me-2-Indan	H	H	5-Bzt	Aa-38	BRA44	C		414 (M <sup>+</sup> +1)
Cb-34	5Me-2-Indan	H	H	1Et-5-1HIdz	Aa-28	BRA44	C		425 (M <sup>+</sup> +1)
Cb-35	4,7DMe-2-Indan	H	H	5-Ind	Aa-2	BRA45	C		410 (M <sup>+</sup> +1)
Cb-36	4,7DMe-2-Indan	H	H	5-1HIdz	Aa-4	BRA45	C		411 (M <sup>+</sup> +1)
Cb-37	4,7DMe-2-Indan	H	H	1Me-5-1HIdz	Aa-5	BRA45	C		425 (M <sup>+</sup> +1)
Cb-38	5,6DMe-2-Indan	H	H	2-Nap	Aa-1	BRA46	C		421 (M <sup>+</sup> +1)
Cb-39	5,6DMe-2-Indan	H	H	1Me-5-1HIdz	Aa-5	BRA46	C		425 (M <sup>+</sup> +1)
Cb-40	5F-2-Indan	H	H	2-Nap	Aa-1	BRA47	C		411 (M <sup>+</sup> +1)
Cb-41	5F-2-Indan	H	H	5-Ind	Aa-2	BRA47	C		400 (M <sup>+</sup> +1)
Cb-42	5F-2-Indan	H	H	5-1HIdz	Aa-4	BRA47	C		401 (M <sup>+</sup> +1)
Cb-43	5F-2-Indan	H	H	1Me-5-1HIdz	Aa-5	BRA47	C		415 (M <sup>+</sup> +1)
Cb-44	5F-2-Indan	H	H	5-Bzt	Aa-38	BRA47	C		418 (M <sup>+</sup> +1)
Cb-45	5F-2-Indan	H	H	3-Qu	Aa-7	BRA47	C		412 (M <sup>+</sup> +1)
Cb-46	5F-2-Indan	H	H	1Et-5-1HIdz	Aa-28	BRA47	C		429 (M <sup>+</sup> +1)
Cb-47	4F-2-Indan	H	H	2-Nap	Aa-1	BRA48	C		411 (M <sup>+</sup> +1)
Cb-48	4F-2-Indan	H	H	1Me-5-Ind	Aa-3	BRA48	C		414 (M <sup>+</sup> +1)
Cb-49	4F-2-Indan	H	H	1Me-5-1HIdz	Aa-5	BRA48	C		415 (M <sup>+</sup> +1)
Cb-50	4,7DF-2-Indan	H	H	2-Nap	Aa-1	BRA49	C		429 (M <sup>+</sup> +1)

Table-Cb-2

Exp.	Rx	Y	Zx	AR	SM1	SM2	LCMS		
							method	RTime	Mass
Cb-51	4,7DF-2-Indan	H	H	1Me-5-Ind	Aa-3	BRA49	C		432 (M <sup>+</sup> +1)
Cb-52	4,7DF-2-Indan	H	H	1Me-5-1HIdz	Aa-5	BRA49	C		433 (M <sup>+</sup> +1)
Cb-53	5,6DF-2-Indan	H	H	2-Nap	Aa-1	BRA50	C		429 (M <sup>+</sup> +1)
Cb-54	5,6DF-2-Indan	H	H	5-Ind	Aa-2	BRA50	C		418 (M <sup>+</sup> +1)
Cb-55	5,6DF-2-Indan	H	H	1Me-5-Ind	Aa-3	BRA50	C		432 (M <sup>+</sup> +1)
Cb-56	5,6DF-2-Indan	H	H	5-1HIdz	Aa-4	BRA50	C		419 (M <sup>+</sup> +1)
Cb-57	5,6DF-2-Indan	H	H	1Me-5-1HIdz	Aa-5	BRA50	C		433 (M <sup>+</sup> +1)
Cb-58	5,6DF-2-Indan	H	H	5-Bzt	Aa-38	BRA50	C		436 (M <sup>+</sup> +1)
Cb-59	5,6DF-2-Indan	H	H	3-Qu	Aa-7	BRA50	C		430 (M <sup>+</sup> +1)
Cb-60	5,6DF-2-Indan	H	H	1Et-5-1HIdz	Aa-28	BRA50	C		447 (M <sup>+</sup> +1)
Cb-61	4Cl-2-Indan	H	H	5-Ind	Aa-2	BRA51	C		416 (M <sup>+</sup> +1)
Cb-62	4Cl-2-Indan	H	H	1Me-5-1HIdz	Aa-5	BRA51	C		431 (M <sup>+</sup> +1)
Cb-63	4Cl-2-Indan	H	H	5-Bzt	Aa-38	BRA51	C		434 (M <sup>+</sup> +1)
Cb-64	5Cl-2-Indan	H	H	2-Nap	Aa-1	BRA52	C		427 (M <sup>+</sup> +1)
Cb-65	5Cl-2-Indan	H	H	5-Ind	Aa-2	BRA52	C		416 (M <sup>+</sup> +1)
Cb-66	5Cl-2-Indan	H	H	1Me-5-1HIdz	Aa-5	BRA52	C		431 (M <sup>+</sup> +1)
Cb-67	5Cl-2-Indan	H	H	3-Qu	Aa-7	BRA52	C		428 (M <sup>+</sup> +1)
Cb-68	5Cl-2-Indan	H	H	1Et-5-1HIdz	Aa-28	BRA52	C		445 (M <sup>+</sup> +1)
Cb-69	4,7DCI-2-Indan	H	H	2-Nap	Aa-1	BRA53	C		462 (M <sup>+</sup> +1)
Cb-70	4,7DCI-2-Indan	H	H	5-Ind	Aa-2	BRA53	C		451 (M <sup>+</sup> +1)
Cb-71	4,7DCI-2-Indan	H	H	1Me-5-1HIdz	Aa-5	BRA53	C		466 (M <sup>+</sup> +1)
Cb-72	5,6DCI-2-Indan	H	H	2-Nap	Aa-1	BRA54	C		462 (M <sup>+</sup> +1)
Cb-73	5,6DCI-2-Indan	H	H	1Me-5-Ind	Aa-3	BRA54	C		465 (M <sup>+</sup> +1)
Cb-74	5,6DCI-2-Indan	H	H	1Me-5-1HIdz	Aa-5	BRA54	C		466 (M <sup>+</sup> +1)
Cb-75	5,6DCI-2-Indan	H	H	5-Bzt	Aa-38	BRA54	C		469 (M <sup>+</sup> +1)
Cb-76	5,6DCI-2-Indan	H	H	3-Qu	Aa-7	BRA54	C		463 (M <sup>+</sup> +1)
Cb-77	5,6DCI-2-Indan	H	H	1Et-5-1HIdz	Aa-28	BRA54	C		480 (M <sup>+</sup> +1)
Cb-78	4MeO-2-Indan	H	H	5-Ind	Aa-2	BRA55	C		412 (M <sup>+</sup> +1)
Cb-79	4MeO-2-Indan	H	H	5-1HIdz	Aa-4	BRA55	C		413 (M <sup>+</sup> +1)
Cb-80	4MeO-2-Indan	H	H	1Me-5-1HIdz	Aa-5	BRA55	C		427 (M <sup>+</sup> +1)
Cb-81	5MeO-2-Indan	H	H	2-Nap	Aa-1	BRA56	C		423 (M <sup>+</sup> +1)
Cb-82	5MeO-2-Indan	H	H	5-Ind	Aa-2	BRA56	C		412 (M <sup>+</sup> +1)
Cb-83	5MeO-2-Indan	H	H	1Me-5-1HIdz	Aa-5	BRA56	C		427 (M <sup>+</sup> +1)
Cb-84	5MeO-2-Indan	H	H	5-Bzt	Aa-38	BRA56	C		430 (M <sup>+</sup> +1)
Cb-90	5,6DMeO-2-Indan	H	H	2-Nap	Aa-1	BRA57	C		453 (M <sup>+</sup> +1)
Cb-91	5,6DMeO-2-Indan	H	H	1Me-5-Ind	Aa-3	BRA57	C		456 (M <sup>+</sup> +1)
Cb-92	5,6DMeO-2-Indan	H	H	1Me-5-1HIdz	Aa-5	BRA57	C		457 (M <sup>+</sup> +1)
Cb-93	5,6DMeO-2-Indan	H	H	5-Bzt	Aa-38	BRA57	C		460 (M <sup>+</sup> +1)
Cb-94	5,6DMeO-2-Indan	H	H	3-Qu	Aa-7	BRA57	C		454 (M <sup>+</sup> +1)
Cb-95	5,6-DMeO-2-Indan	H	H	1Et-5-1HIdz	Aa-28	BRA57	C		471 (M <sup>+</sup> +1)
Cb-85	cHex	H	H	2-Nap	Aa-1	BRA58	C		359 (M <sup>+</sup> +1)
Cb-86	cHex	H	H	5-Ind	Aa-2	BRA58	C		348 (M <sup>+</sup> +1)
Cb-87	cHex	H	H	1Me-5-Ind	Aa-3	BRA58	C		362 (M <sup>+</sup> +1)
Cb-88	cHex	H	H	5-1HIdz	Aa-4	BRA58	C		349 (M <sup>+</sup> +1)
Cb-89	cHex	H	H	1Me-5-1HIdz	Aa-5	BRA58	C		363 (M <sup>+</sup> +1)

[Reference Examples: Intermediate Ab-1 to Ab-47]

Synthesis of methyl 3-[4-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)-3-(naphthalen-2-yl)phenyl]propionate (Intermediate Ab-1)

Compound No. Aa-1 (253.2 mg), bispinacolate diboron (202.6mg, Ald), PdCl<sub>2</sub>(dppf) (43.4 mg) and potassium acetate (289 mg) were added to DMF (5.7 ml),

and stirred with heating at 80°C for 20 hours under argon gas atmosphere. The reaction mixture was added with ethyl acetate (200 ml), washed with saturated brine, and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 4:1) to obtain the title compound (Intermediate Ab-1, 194.6 mg).

Typical examples of the compounds of the present invention including those mentioned above that can be obtained by reacting and treating corresponding starting compounds according to the synthesis method of Intermediate Ab-1 are shown in Table-Ab-1.

In the column indicated as "Mass" in the table, data of mass spectra measured by fast atom bombardment mass spectrometry (FAB-MS) are shown.

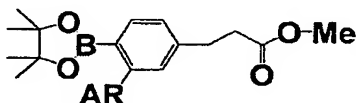


Table-Ab-1

Exp.	AR	Mass	Exp.	AR	Mass
Ab-1	2-Nap	417 ( $M^+ + 1$ )	Ab-25	1Me-4-1HIdz	421 ( $M^+ + 1$ )
Ab-2	5-Ind	406 ( $M^+ + 1$ )	Ab-26	5-1HIdz	407 ( $M^+ + 1$ )
Ab-3	1Me-5-Ind	420 ( $M^+ + 1$ )	Ab-27	1Me-5-1HIdz	421 ( $M^+ + 1$ )
Ab-4	5-1HIdz	407 ( $M^+ + 1$ )	Ab-28	1Et-5-1HIdz	435 ( $M^+ + 1$ )
Ab-5	1Me-5-1HIdz	421 ( $M^+ + 1$ )	Ab-29	1Pr-5-1HIdz	449 ( $M^+ + 1$ )
Ab-6	5-BF	410 ( $M^+ + 1$ )	Ab-30	2Me-5-2HIdz	421 ( $M^+ + 1$ )
Ab-7	3-Qu	418 ( $M^+ + 1$ )	Ab-31	6-1HIdz	407 ( $M^+ + 1$ )
Ab-8	1-Nap	417 ( $M^+ + 1$ )	Ab-32	1Me-6-1HIdz	421 ( $M^+ + 1$ )
Ab-9	6MeO-2-Nap	447 ( $M^+ + 1$ )	Ab-33	3Me-5-1HIdz	421 ( $M^+ + 1$ )
Ab-10	6(Me <sub>2</sub> N)-2-Nap	460 ( $M^+ + 1$ )	Ab-34	1,3DMe-5-1HIdz	435 ( $M^+ + 1$ )
Ab-11	4-Ind	406 ( $M^+ + 1$ )	Ab-35	5-BT	423 ( $M^+ + 1$ )
Ab-12	1Me-4-Ind	420 ( $M^+ + 1$ )	Ab-36	2,3DMe-5-BF	435 ( $M^+ + 1$ )
Ab-13	6-Ind	406 ( $M^+ + 1$ )	Ab-37	5-2ABzt	439 ( $M^+ + 1$ )
Ab-14	1Me-6-Ind	420 ( $M^+ + 1$ )	Ab-38	5-Bzt	434 ( $M^+ + 1$ )
Ab-15	2-Ind	406 ( $M^+ + 1$ )	Ab-39	2Me-5-Bzt	438 ( $M^+ + 1$ )
Ab-16	1Me-2-Ind	420 ( $M^+ + 1$ )	Ab-40	2,2DMe-5-2ABzt	467 ( $M^+ + 1$ )
Ab-17	3-Ind	406 ( $M^+ + 1$ )	Ab-41	6-2ABzt	439 ( $M^+ + 1$ )
Ab-18	1Me-3-Ind	420 ( $M^+ + 1$ )	Ab-42	6-Bzt	434 ( $M^+ + 1$ )
Ab-19	1iPr-5-Ind	448 ( $M^+ + 1$ )	Ab-43	2Me-6-Bzt	438 ( $M^+ + 1$ )
Ab-20	1cPen-5-Ind	474 ( $M^+ + 1$ )	Ab-44	6-Qu	418 ( $M^+ + 1$ )
Ab-21	3Me-5-Ind	420 ( $M^+ + 1$ )	Ab-45	6-1Q	418 ( $M^+ + 1$ )
Ab-22	1,3DMe-5Ind	434 ( $M^+ + 1$ )	Ab-46	2-BF	407 ( $M^+ + 1$ )
Ab-23	1,2,3triMe-5Ind	448 ( $M^+ + 1$ )	Ab-47	2-BT	423 ( $M^+ + 1$ )
Ab-24	4-1HIdz	407 ( $M^+ + 1$ )			

[Example Da-1]



Synthesis of methyl 3-[4-(phenylmethyl)-3-(naphthalen-2-yl)phenyl]propionate  
(Compound No. Da-1)

According to a procedure described in literature (S. Chowdhury et al., Tetrahedron. Lett., 1999, p.7599), (Ph<sub>3</sub>P)<sub>4</sub>Pd (14.8 mg) and a solution of benzyl bromide (corresponding to the substance mentioned in the column of SM2 in Table Da-1 mentioned later) in dimethoxyethane (1.3 ml) were stirred with heating at 50°C for 10 minutes under argon atmosphere, then added with Compound Ab-1 (52.4 mg, corresponding to the substance mentioned in the column of SM1 in Table Da-1 mentioned later), and 2 N sodium carbonate (160  $\mu$ l), and refluxed by heating for 58 hours. The reaction mixture was added with ethyl acetate (60 ml), washed successively with saturated aqueous sodium hydrogencarbonate and saturated brine, dried, and then concentrated under reduced pressure. The residue was purified by flash column chromatography (hexane:ethyl acetate = 8:1) to obtain the title compound (Compound No. Da-1, 33.2 mg).

[Example Da-2]

Synthesis of 3-[4-(phenylmethyl)-3-(naphthalen-2-yl)phenyl]propionic acid  
(Compound No. Da-2)

According to the procedure described in the synthesis method of Compound Ca-2 provided that the reaction was performed for 3 hours, Compound No. Da-1 (28.2 mg) and 2 N aqueous sodium hydroxide (38  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. Da-2, 23.7 mg).

[Examples Da-1 to Da-70]

Typical examples of the compounds of the present invention including those mentioned in the examples described above, that can be obtained by reacting and treating corresponding starting compounds according to the methods described in Examples Da-1 and Da-2, are shown in Table Da-1 and Table Da-2.

The substances mentioned in the columns of "SM1" in the tables correspond

to reaction intermediates, and those mentioned in the columns of "SM2" in the tables correspond to the acid halide mentioned in Example Da-1. The halide reagents mentioned in the columns of "SM2" with the symbols of "HAL (number))" are those mentioned in Table-Ha. The reagents for which cells of the columns of "Manufacturer" are blank in the tables are synthesized according to a method described in ordinary chemical literature.

Table-Ha

Reagent	Name of reagent	Manufacturer
HAL-1	Benzyl bromide	Ald
HAL-2	4-Methoxybenzyl bromide	Ald
HAL-3	3-Methoxybenzyl bromide	Ald
HAL-4	2-Methoxybenzyl bromide	Ald
HAL-5	4-Methylbenzyl bromide	Ald
HAL-6	3-Methylbenzyl bromide	Ald
HAL-7	2-Methylbenzyl bromide	Ald
HAL-8	4-Trifluoromethylbenzyl bromide	Ald
HAL-9	3-Trifluoromethylbenzyl bromide	Ald
HAL-10	2-Trifluoromethylbenzyl bromide	Ald
HAL-11	4-Chlorobenzyl bromide	Ald
HAL-12	3-Chlorobenzyl bromide	Ald
HAL-13	2-Chlorobenzyl bromide	Ald
HAL-14	4-Fluorobenzyl bromide	Ald
HAL-15	3-Fluorobenzyl bromide	Ald
HAL-16	2-Fluorobenzyl bromide	Ald
HAL-17	1-Bromo-2-phenyl ethane	Ald
HAL-18	1-Bromo-2-(4-chloro phenyl) ethane	Ald
HAL-19	1-Bromo-2-(3-chloro phenyl) ethane	
HAL-20	1-Bromo-2-(2-chloro phenyl) ethane	
HAL-21	1-Bromo-2-(4-dimethyl aminophenyl) ethane	
HAL-22	Benzoyl chloride	TCI
HAL-23	Acetyl chloride	WAKO
HAL-24	i-Butyryl chloride	Ald
HAL-25	Cyclohexylcarbonyl chloride	Ald
HAL-26	4-Methoxybenzoyl chloride	TCI
HAL-27	4-Methylbenzoyl chloride	Ald
HAL-28	4-Chlorobenzoyl chloride	TCI
HAL-29	Phenylacetyl chloride	WAKO
HAL-30	2-Phenylpropionyl chloride	TCI

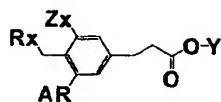


Table-Da-1

Exp.	Rx	Y	Zx	AR	SM1	SM2	LCMS		
							method	RTTime	Mass
Da-1	Ph	Me	H	2-Nap	Ab-1	HAL-1	D		N.D
Da-2	Ph	H	H	2-Nap	Da-1	-	C		367 (M <sup>+</sup> +1)
Da-3	Ph	Me	H	5-Ind	Ab-2	HAL-1			
Da-4	Ph	H	H	5-Ind	Da-3	-			
Da-5	Ph	Me	H	1Me-5-Ind	Ab-3	Ha-1	C		384 (M <sup>+</sup> +1)
Da-6	Ph	H	H	1Me-5-Ind	Da-5	-	C		369 (M <sup>+</sup> +1)
Da-7	Ph	Me	H	5-1HIdz	Ab-4	Ha-1			
Da-8	Ph	H	H	5-1HIdz	Da-7	-			
Da-9	Ph	Me	H	1Me-5-1HIdz	Ab-5	HAL-1	C		385 (M <sup>+</sup> +1)
Da-10	Ph	H	H	1Me-5-1HIdz	Da-9	-	C		370 (M <sup>+</sup> +1)
Da-11	4MeOPh	H	H	2-Nap	Ab-1	HAL-2			
Da-12	4MeOPh	H	H	5-Ind	Ab-2	HAL-2			
Da-13	4MeOPh	H	H	1Me-5-1HIdz	Ab-5	HAL-2			
Da-14	3MeOPh	H	H	2-Nap	Ab-1	HAL-3	C		397 (M <sup>+</sup> +1)
Da-15	3MeOPh	H	H	5-Ind	Ab-2	HAL-3			
Da-16	3MeOPh	H	H	1Me-5-1HIdz	Ab-5	HAL-3			
Da-17	2MeOPh	H	H	2-Nap	Ab-1	HAL-4			
Da-18	2MeOPh	H	H	5-Ind	Ab-2	HAL-4			
Da-19	2MeOPh	H	H	1Me-5-1HIdz	Ab-5	HAL-4			
Da-20	4MePh	H	H	2-Nap	Ab-1	HAL-5	C		381 (M <sup>+</sup> +1)
Da-21	4MePh	H	H	5-Ind	Ab-2	HAL-5			
Da-22	4MePh	H	H	1Me-5-1HIdz	Ab-5	HAL-5			
Da-23	3MePh	H	H	2-Nap	Ab-1	HAL-6			
Da-24	3MePh	H	H	5-Ind	Ab-2	HAL-6			
Da-25	3MePh	H	H	1Me-5-1HIdz	Ab-5	HAL-6			
Da-26	2MePh	H	H	2-Nap	Ab-1	HAL-7	C		381 (M <sup>+</sup> +1)
Da-27	2MePh	H	H	5-Ind	Ab-2	HAL-7	C		370 (M <sup>+</sup> +1)
Da-28	2MePh	H	H	1Me-5-1HIdz	Ab-5	HAL-7			
Da-29	4CF <sub>3</sub> Ph	H	H	2-Nap	Ab-1	HAL-8			
Da-30	4CF <sub>3</sub> Ph	H	H	5-Ind	Ab-2	HAL-8			
Da-31	4CF <sub>3</sub> Ph	H	H	1Me-5-1HIdz	Ab-5	HAL-8			
Da-32	3CF <sub>3</sub> Ph	H	H	2-Nap	Ab-1	HAL-9			
Da-33	3CF <sub>3</sub> Ph	H	H	5-Ind	Ab-2	HAL-9			
Da-34	3CF <sub>3</sub> Ph	H	H	1Me-5-1HIdz	Ab-5	HAL-9			
Da-35	2CF <sub>3</sub> Ph	H	H	2-Nap	Ab-1	HAL-10			
Da-36	2CF <sub>3</sub> Ph	H	H	5-Ind	Ab-2	HAL-10			
Da-37	2CF <sub>3</sub> Ph	H	H	1Me-5-1HIdz	Ab-5	HAL-10			
Da-38	4ClPh	H	H	2-Nap	Ab-1	HAL-11	C		401 (M <sup>+</sup> +1)
Da-39	4ClPh	H	H	5-Ind	Ab-2	HAL-11	C		390 (M <sup>+</sup> +1)
Da-40	4ClPh	H	H	1Me-5-1HIdz	Ab-5	HAL-11			
Da-41	3ClPh	H	H	2-Nap	Ab-1	HAL-12			
Da-42	3ClPh	H	H	5-Ind	Ab-2	HAL-12			
Da-43	3ClPh	H	H	1Me-5-1HIdz	Ab-5	HAL-12			
Da-44	2ClPh	H	H	2-Nap	Ab-1	HAL-13			
Da-45	2ClPh	H	H	5-Ind	Ab-2	HAL-13			
Da-46	2ClPh	H	H	1Me-5-1HIdz	Ab-5	HAL-13			
Da-47	4FPh	H	H	2-Nap	Ab-1	HAL-14	C		385 (M <sup>+</sup> +1)
Da-48	4FPh	H	H	5-Ind	Ab-2	HAL-14			
Da-49	4FPh	H	H	1Me-5-1HIdz	Ab-5	HAL-14	C		389 (M <sup>+</sup> +1)
Da-50	3FPh	H	H	2-Nap	Ab-1	HAL-15			

Table-Da-2

Exp.	Rx	Y	Zx	AR	SM1	SM2	LCMS		
							method	RTime	Mass
Da-51	3FPh	H	H	5-Ind	Ab-2	HAL-15			
Da-52	3FPh	H	H	1Me-5-1HIdz	Ab-5	HAL-15			
Da-53	2FPh	H	H	2-Nap	Ab-1	HAL-16			
Da-54	2FPh	H	H	5-Ind	Ab-2	HAL-16			
Da-55	2FPh	H	H	1Me-5-1HIdz	Ab-5	HAL-16			
Da-56	Bn	H	H	2-Nap	Ab-1	HAL-17	C		381 (M <sup>+</sup> +1)
Da-57	Bn	H	H	5-Ind	Ab-2	HAL-17			
Da-58	Bn	H	H	1Me-5-1HIdz	Ab-5	HAL-17	C		419 (M <sup>+</sup> +1)
Da-59	4ClBn	H	H	2-Nap	Ab-1	HAL-18			
Da-60	4ClBn	H	H	5-Ind	Ab-2	HAL-18			
Da-61	4ClBn	H	H	1Me-5-1HIdz	Ab-5	HAL-18	C		385 (M <sup>+</sup> +1)
Da-62	3ClBn	H	H	2-Nap	Ab-1	HAL-19	C		415 (M <sup>+</sup> +1)
Da-63	3ClBn	H	H	5-Ind	Ab-2	HAL-19			
Da-64	3ClBn	H	H	1Me-5-1HIdz	Ab-5	HAL-19			
Da-65	2ClBn	H	H	2-Nap	Ab-1	HAL-20			
Da-66	2ClBn	H	H	5-Ind	Ab-2	HAL-20			
Da-67	2ClBn	H	H	1Me-5-1HIdz	Ab-5	HAL-20			
Da-68	4DMABn	H	H	2-Nap	Ab-1	HAL-21	C		424 (M <sup>+</sup> +1)
Da-69	4DMABn	H	H	5-Ind	Ab-2	HAL-21	C		413 (M <sup>+</sup> +1)
Da-70	4DMABn	H	H	1Me-5-1HIdz	Ab-5	HAL-21			

## [Example Ea-1]

Synthesis of methyl 3-[4-(phenylcarbonyl)-3-(naphthalen-2-yl)phenyl]propionate  
(Compound No. Ea-1)

According to a procedure described in literature (Y. Urawa et al, Tetrahedron. Lett., 2003, p.271), Compound Ab-1 (112.1mg, corresponding to the substance mentioned in the column of SM1 in Table-Ea-1 mentioned later), dichlorobis(triphenylphosphine)palladium (18.9 mg, KANTO), and a solution of potassium phosphate (147.1 mg) in toluene (2.6 ml) were added with benzoyl chloride (47  $\mu$ g, corresponding to the substance mentioned in the column of SM2 in Table-Ea-1), and stirred with heating at 110°C for 48 hours under nitrogen atmosphere. The reaction mixture was washed successively with saturated aqueous sodium hydrogencarbonate, water and saturated brine, dried, and then concentrated under reduced pressure. The residue was purified by flash column chromatography (hexane:ethyl acetate = 7:1) to obtain the title compound (Compound No. Ea-1, 88.3 mg).

## [Example Ea-2]

Synthesis of 3-[4-phenylcarbonyl-3-(naphthalen-2-yl)phenyl]propionic acid  
(Compound No. Ea-2)

According to the procedure described in the synthesis method of Compound Ca-2 with the modification that the reaction was carried out for 3 hour, Compound No. Ea-1 (82.6 mg) and 2 N aqueous sodium hydroxide (105 ml) were reacted and treated to obtain the title compound (Compound No. Ea-2, 70.7 mg).

[Examples Ea-1 to Ea-34]

Typical examples of the compounds of the present invention including those mentioned in the examples described above, that can be obtained by reacting and treating corresponding starting compounds according to the methods described in Examples Ea-1 and Ea-2, are shown in Table-Ea-1.

The substances mentioned in the column of "SM1" in the table correspond to reaction intermediates, and those mentioned in the column of "SM2" in the table correspond to acid chlorides mentioned in Table Ea-1. The acid chlorides mentioned with the symbols of "HAL (number)" in the column of "SM2" are those mentioned in Table-Ha.

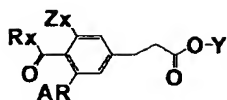


Table-Ea-1

Exp.	Rx	Y	Zx	AR	SM1	SM2	LCMS		
							method	RTime	Mass
Ea-1	Ph	Me	H	2-Nap	Ab-1	HAL-22	C		395 (M <sup>+</sup> +1)
Ea-2	Ph	H	H	2-Nap	Ea-1	-	C		381 (M <sup>+</sup> +1)
Ea-3	Ph	Me	H	5-Ind	Ab-2	HAL-22			
Ea-4	Ph	H	H	5-Ind	Ea-3	-			
Ea-5	Ph	Me	H	1Me-5-Ind	Ab-3	HAL-22	C		398 (M <sup>+</sup> +1)
Ea-6	Ph	H	H	1Me-5-Ind	Ea-5	-	C		384 (M <sup>+</sup> +1)
Ea-7	Ph	Me	H	5-1HIdz	Ab-4	HAL-22			
Ea-8	Ph	H	H	5-1HIdz	Ea-7	-			
Ea-9	Ph	Me	H	1Me-5-1HIdz	Ab-5	HAL-22	C		399 (M <sup>+</sup> +1)
Ea-10	Ph	H	H	1Me-5-1HIdz	Ea-9	-	C		385 (M <sup>+</sup> +1)
Ea-11	Me	H	H	2-Nap	Ab-1	HAL-23	C		319 (M <sup>+</sup> +1)
Ea-12	Me	H	H	5-Ind	Ab-2	HAL-23	C		308 (M <sup>+</sup> +1)
Ea-13	Me	H	H	1Me-5-1HIdz	Ab-5	HAL-23			
Ea-14	iBu	H	H	2-Nap	Ab-1	HAL-24	C		361 (M <sup>+</sup> +1)
Ea-15	iBu	H	H	5-Ind	Ab-2	HAL-24			
Ea-16	iBu	H	H	1Me-5-1HIdz	Ab-5	HAL-24	C		365 (M <sup>+</sup> +1)
Ea-17	cHex	H	H	2-Nap	Ab-1	HAL-25	C		386 (M <sup>+</sup> +1)
Ea-18	cHex	H	H	5-Ind	Ab-2	HAL-25			
Ea-19	cHex	H	H	1Me-5-1HIdz	Ab-5	HAL-25			
Ea-20	4MeOPh	H	H	2-Nap	Ab-1	HAL-26	C		411 (M <sup>+</sup> +1)
Ea-21	4MeOPh	H	H	5-Ind	Ab-2	HAL-26	C		400 (M <sup>+</sup> +1)
Ea-22	4MeOPh	H	H	1Me-5-1HIdz	Ab-5	HAL-26			
Ea-23	4MePh	H	H	2-Nap	Ab-1	HAL-27			
Ea-24	4MePh	H	H	5-Ind	Ab-2	HAL-27			
Ea-25	4MePh	H	H	1Me-5-1HIdz	Ab-5	HAL-27			
Ea-26	4ClPh	H	H	2-Nap	Ab-1	HAL-28	C		415 (M <sup>+</sup> +1)
Ea-27	4ClPh	H	H	5-Ind	Ab-2	HAL-28			
Ea-28	4ClPh	H	H	1Me-5-1HIdz	Ab-5	HAL-28			
Ea-29	Bn	H	H	2-Nap	Ab-1	HAL-29	C		395 (M <sup>+</sup> +1)
Ea-30	Bn	H	H	5-Ind	Ab-2	HAL-29			
Ea-31	Bn	H	H	1Me-5-1HIdz	Ab-5	HAL-29	C		399 (M <sup>+</sup> +1)
Ea-32	1PhEt	H	H	2-Nap	Ab-1	HAL-30	C		409 (M <sup>+</sup> +1)
Ea-33	1PhEt	H	H	5-Ind	Ab-2	HAL-30	C		398 (M <sup>+</sup> +1)
Ea-34	1PhEt	H	H	1Me-5-1HIdz	Ab-5	HAL-30			

[Reference Examples: Intermediate Ac-1 and Ac-2]

Synthesis of t-butyltrimethylsilyl 3-[3-bromo-4-(t-butyltrimethylsilyloxy)phenyl]acrylate (Intermediate Ac-1)

According to the procedure described in the synthesis method of Intermediate 43, 3-[3-bromo-4-hydroxyphenyl]acrylic acid (12.01 g) obtainable from 4-hydroxybenzaldehyde (TCI) by a method known from literature (Y. Nagao et al., Tetrahedron Lett., 1980, p.4931) was reacted with imidazole (16.01 g) and t-butyltrimethylsilyl chloride (7.43 g) and treated to obtain the title compound (Intermediate Ac-1, 17.43 g).

Synthesis of 3-[3-bromo-4-(t-butyltrimethylsilyloxy)phenyl]acrylic acid

(Intermediate Ac-2)

A solution of Compound Ac-1 (17.43 g) in methanol (100 ml) was added with 1 N hydrochloric acid (5 ml), and stirred at room temperature for 3 hours. The reaction solution was extracted with ethyl acetate (500 ml), and washed with saturated brine. The organic layer was dried, and then the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography (hexane:ethyl acetate = 6:1) to obtain the title compound (Compound No. Ac-2, 14.60 g).

[Example Ga-1]

Synthesis of methyl 3-[3-(1H-indol-5-yl)-4-(3-pyridinemethyloxy)phenyl]acrylate  
(Compound No. Ga-1)

(Step 1)

A solution of Compound Ac-2 (3.06 g), diisopropyl carbodiimide (henceforth abbreviated as "DIC", 1.33 ml) and dimethylaminopyridine (86.8 mg) in DMF (100 ml) was added with SynPhase-PS-D-series Lantern, Hydroxymethylphenoxy Linker (henceforth abbreviated as "PSL", 0.035 mmol per lantern, 81 lanterns, Mimotopes), and left standing at room temperature for 16 hours. After the reaction mixture was removed, PSL was washed successively with DMF (100 ml), methanol (100 ml), dichloromethane (100 ml), and THF (100 ml) three times for each, and dried under reduced pressure.

PSL (81 lanterns mentioned above) was added to a solution of tetrabutylammonium fluoride (8.5 ml, Ald, 1 N THF solution) in THF (80 ml), and left standing at room temperature for 23 hours. After the reaction mixture was removed, PSL was successively washed with DMF (100 ml) three times, alternately with DMF:water:acetic acid (75:25:1, 100 ml) and methanol:water:acetic acid (75:25:1, 100 ml) twice for each, alternately with DMF:water (4:1, 100 ml) and methanol:water (4:1, 100 ml) twice for each, and with THF (100 ml), chloroform



(100 ml), DMF (100 ml), and chloroform (100 ml) twice for each, and then dried under reduced pressure to obtain PLS-1 (81 lanterns).

(Step 2)

PSL-1 (3 lanterns out of those mentioned above) was added to a mixed solution of 3-pyridinemethanol (147.6  $\mu$ l, corresponding to the substance mentioned in the column of SM1 in Table-Ga-1 mentioned later), DBAB (242.1mg, Sigma) and  $\text{Ph}_3\text{P}$  (275.6 mg, KANTO) in dehydrated THF (3.24 ml), and left standing at room temperature for 15 hours. After the reaction mixture was removed, PSL was successively washed with THF (3.5 ml) and DMF (3.5 ml) four times for each, alternately with methanol (3.5 ml) and DMF (3.5 ml) twice for each, alternately with DMF (3.5 ml) and dichloromethane (3.5 ml) twice for each, with dichloromethane (3.5 ml) twice, and dried under reduced pressure to obtain PSL-2 (3 vials).

(Step 3)

PSL-2 (1 lantern out of those mentioned above) was added to a mixed solution of 1H-indole-5-boronic acid (11.3 mg, corresponding to the substance mentioned in the column of SM2 in Table-Ga-1 mentioned later),  $(\text{Ph}_3\text{P})_4\text{Pd}$  (8.1 mg), and 2 N aqueous cesium carbonate (176  $\mu$ l) in DMF (800  $\mu$ l), and heated at 80°C for 18 hours under argon atmosphere. After the reaction mixture was removed, PSL was successively washed with DMF (1.0 ml) four times, with methanol (1.0 ml) twice, alternately with DMF (1.0 ml) and methanol (1.0 ml) twice for each, alternately with DMF (1.0 ml) and dichloromethane (1.0 ml) twice for each, and with dichloromethane (1.0 ml) twice, and dried under reduced pressure. This PSL was added to a solution of sodium methoxide (175  $\mu$ l, WAKO, 1 N solution in methanol) in THF:methanol (2:1, 1.5 ml), and left standing at room temperature for 19 hours. After the reaction, PSL was removed, and the reaction solution was added with water (500  $\mu$ l), and stirred with heating at 60°C for 3 hours. The

reaction solution was concentrated under reduced pressure, then added with water (200  $\mu$ l) and chloroform (1 ml), and passed through a diatomaceous earth column, and the obtained filtrate was concentrated under reduced pressure to obtain the title compound (Compound No. Ga-1, 10.6 mg).

[Examples Ga-1 to Ga-55]

Typical examples of the compounds of the present invention including those mentioned in the examples described above, that can be obtained by reacting and treating corresponding starting compounds according to the method described in Example Ga-1, are shown in Table-Ga-1 and Table-Ga-2.

The substances mentioned in the columns of "SM1" in the tables correspond to the alcohol reagent mentioned in Example Ga-1, and those mentioned in the columns of "SM2" in the tables correspond to the boronic acid reagent mentioned in Table Ga-1. The alcohol reagents mentioned in the columns of "SM1" with the symbols of "ALC (number))" are those mentioned in Table-I. The boronic acid reagents mentioned with the symbols of "BRA (number))" in the columns of "SM2" are those mentioned in Table-Ba-1 and Table-Ba-2.

Table-I

Reagent	Name of reagent	Manufacture	Reagent	Name of reagent	Manufacture
ALC-1	Cyclopentanol	KANTO	ALC-16	2-Phenylthio ethanol	TCI
ALC-2	Cyclohexanol	Ald	ALC-17	5-(2-Hydroxyethyl)- 4-methylthiazol	TCI
ALC-3	Benzyl Alcohol	Ald	ALC-18	1-Butanol	TCI
ALC-4	2-Methyl-1-propyl alcohol	TCI	ALC-19	2-Hydroxyethyl acetate	TCI
ALC-5	4-Fluorophenetyl alcohol	Ald	ALC-20	N-(2-Hydroxyethyl) morpholine	TCI
ALC-6	1-Phenylethanol	WAKO	ALC-21	2-(2- Dimethylaminoethoxy)	TCI
ALC-7	2-(N-Methylanilino) ethanol	TCI	ALC-22	Methyl glycolate	TCI
ALC-8	2-Hydroxy indane	TCI	ALC-23	1-Phenyl ethanol	TCI
ALC-9	2-Hydroxymethyl- 1,4-benzodioxane	TCI	ALC-24	2-Chlorobenzyl alcohol	TCI
ALC-10	2-(4-Dimethyl) phenyl ethanol	Ald	ALC-25	3-Chlorobenzyl alcohol	TCI
ALC-11	3-Pyridine methanol	TCI	ALC-26	4-Chlorobenzyl alcohol	TCI
ALC-12	m-Chlorobenzyl alcohol	TCI	ALC-27	2-Methoxybenzyl alcohol	TCI
ALC-13	4-n-Butoxybenzyl alcohol	TCI	ALC-28	3-Methoxybenzyl alcohol	TCI
ALC-14	2-Hydroxyacetophenone	TCI	ALC-29	4-Methoxybenzyl alcohol	TCI
ALC-15	2-Phenoxy ethanol	TCI			

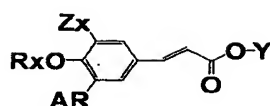


Table-Ga-1

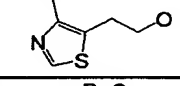
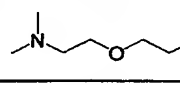
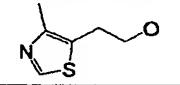
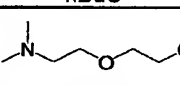
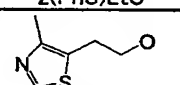
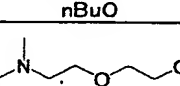
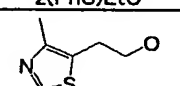
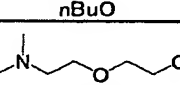
Exp.	RxO	Y	Zx	AR	SM1	SM2	LCMS		
							method	RTime	Mass
Ga-1	3PyMeO	H	H	5-Ind	ALC-11	BRA2	A	3.27	371 (M <sup>+</sup> +1)
Ga-2	2(PhS)EtO	H	H	5-Ind	ALC-16	BRA2			
Ga-3		H	H	5-Ind	ALC-17	BRA2	A	3.07	405 (M <sup>+</sup> +1)
Ga-4	nBuO	H	H	5-Ind	ALC-18	BRA2			
Ga-5		H	H	5-Ind	ALC-21	BRA2			
Ga-6	cPenO	H	H	5-Ind	ALC-1	BRA2			
Ga-7	cHexO	H	H	5-Ind	ALC-2	BRA2			
Ga-8	PhMeO	H	H	5-Ind	ALC-3	BRA2	A	3.79	356 (M <sup>+</sup> +1)
Ga-9	cPenO	H	H	2-BF	ALC-1	BRA18			
Ga-10	cHexO	H	H	2-BF	ALC-2	BRA18			
Ga-11	2-IndanO	H	H	2-BF	ALC-8	BRA18	A	3.85	397 (M <sup>+</sup> +1)
Ga-12	3PyMeO	H	H	2-BF	ALC-11	BRA18			
Ga-13	2(PhS)EtO	H	H	2-BF	ALC-16	BRA18	A	3.61	417 (M <sup>+</sup> +1)
Ga-14		H	H	2-BF	ALC-17	BRA18			
Ga-15	nBuO	H	H	2-BF	ALC-18	BRA18			
Ga-16		H	H	2-BF	ALC-21	BRA18			
Ga-17	cPenO	H	H	1Me-5-1HIdz	ALC-1	BRA6			
Ga-18	cHexO	H	H	1Me-5-1HIdz	ALC-2	BRA6	A	3.74	377 (M <sup>+</sup> +1)
Ga-19	2-IndanO	H	H	1Me-5-1HIdz	ALC-8	BRA6			
Ga-20	3PyMeO	H	H	1Me-5-1HIdz	ALC-11	BRA6			
Ga-21	2(PhS)EtO	H	H	1Me-5-1HIdz	ALC-16	BRA6			
Ga-22		H	H	1Me-5-1HIdz	ALC-17	BRA6			
Ga-23	nBuO	H	H	1Me-5-1HIdz	ALC-18	BRA6			
Ga-24		H	H	1Me-5-1HIdz	ALC-21	BRA6			
Ga-25	3PyMeO	H	H	1-Nap	ALC-11	BRA16			
Ga-26	2(PhS)EtO	H	H	1-Nap	ALC-16	BRA16	C		427 (M <sup>+</sup> +1)
Ga-27		H	H	1-Nap	ALC-17	BRA16			
Ga-28	nBuO	H	H	1-Nap	ALC-18	BRA16			
Ga-29		H	H	1-Nap	ALC-21	BRA16			
Ga-30	1PhEtO	H	H	5-Ind	ALC-6	BRA2			

Table-Ga-2

Exp.	RxO	Y	Zx	AR	SM1	SM2	LCMS		
							method	RTTime	Mass
Ga-31	1PhEtO	H	H	2-BF	ALC-6	BRA18			
Ga-32	1PhEtO	H	H	1Me-5-1Hidz	ALC-6	BRA6	A	3.55	399 (M <sup>+</sup> +1)
Ga-33	1PhEtO	H	H	1-Nap	ALC-6	BRA16			
Ga-34	1PhEtO	H	H	2-Nap	ALC-6	BRA1			
Ga-35	1PhEtO	H	H	2-Nap	ALC-6	BRA1	C		395 (M <sup>+</sup> +1)
Ga-36	1PhEtO	H	H	5-Ind	ALC-6	BRA2			
Ga-37	2ClPhMeO	H	H	2-Nap	ALC-24	BRA1			
Ga-38	2ClPhMeO	H	H	5-Ind	ALC-24	BRA2			
Ga-39	3ClPhMeO	H	H	2-Nap	ALC-25	BRA1	C		415 (M <sup>+</sup> +1)
Ga-40	3ClPhMeO	H	H	5-Ind	ALC-25	BRA2			
Ga-41	4ClPhMeO	H	H	2-Nap	ALC-26	BRA1			
Ga-42	4ClPhMeO	H	H	5-Ind	ALC-26	BRA2	C		404 (M <sup>+</sup> +1)
Ga-43	2MeOPhMeO	H	H	2-Nap	ALC-27	BRA1			
Ga-44	2MeOPhMeO	H	H	5-Ind	ALC-27	BRA2			
Ga-45	3MeOPhMeO	H	H	2-Nap	ALC-28	BRA1			
Ga-46	3MeOPhMeO	H	H	5-Ind	ALC-28	BRA2			
Ga-47	4MeOPhMeO	H	H	2-Nap	ALC-29	BRA1			
Ga-48	4MeOPhMeO	H	H	5-Ind	ALC-29	BRA2			
Ga-49	nBuO	H	H	3-Qu	ALC-18	BRA10	C		348 (M <sup>+</sup> +1)
Ga-50	nBuO	H	H	3-Thienyl	ALC-18	BRA36			
Ga-51	nBuO	H	H	4-Py	ALC-18	BRA26			
Ga-52	nBuO	H	H	cPen	ALC-18	BRA28			
Ga-53	nBuO	H	H	2FPh	ALC-18	BRA32	C		315 (M <sup>+</sup> +1)
Ga-54	nBuO	H	H	3FPh	ALC-18	BRA33			
Ga-55	nBuO	H	H	4FPh	ALC-18	BRA34			

[Reference Examples: Intermediate s-1 to s-52]

Synthesis of methyl 3-[4-(4-methylphenylthio)-3-nitrophenyl]acrylate (Intermediate s-1) (Synthesis method SF)

A solution of 3-[4-(4-methylphenylthio)-3-nitrophenyl]acrylic acid (631 mg, MAYB) in a mixture of methanol (12.6 ml), ethyl acetate (6.3 ml) and THF (6.3 ml) was added dropwise to methanol (12.6 ml) beforehand under ice cooling, and then the mixture was added with a solution of thionyl chloride (735  $\mu$ l, KANTO) in methanol (50 ml) under ice cooling, stirred for 30 minutes, then warmed to room temperature, and further stirred for 15.5 hours. The reaction mixture was poured into aqueous sodium hydrogencarbonate (50 ml) for neutralization, and extracted with ethyl acetate (50 ml), and the organic layer was washed with saturated brine. The organic layer was dried, and then the solvent was evaporated under reduced pressure to obtain the title compound (Intermediate s-1, 659 mg).

Synthesis of methyl 3-[4-(4-methylphenylthio)-3-nitrophenyl]propionate

(Intermediate s-2) (Synthesis method SD1)

A solution of Intermediate s-1 (494 mg) in ethyl acetate (75 ml) was added with 10% palladium hydroxide/carbon (150 mg, NE CHEMCAT), and stirred at room temperature for 14 hours under hydrogen atmosphere. The reaction mixture was filtered, and the solvent of the filtrate was evaporated under reduced pressure. The residue was dissolved in methanol (75 ml) again, added with 5 N hydrochloric acid (600  $\mu$ l) and 10% palladium hydroxide/carbon (151 mg), and stirred at room temperature for 22 hours under hydrogen atmosphere. The reaction mixture was filtered, and the solvent of the filtrate was evaporated under reduced pressure to obtain the title compound (Intermediate s-2, 419 mg).

Synthesis of methyl 3-[3-bromo-4-(4-methylphenylthio)phenyl]propionate

(Intermediate s-3) (Synthesis method SE1)

A solution of hydrobromic acid (690  $\mu$ l) in methanol (3.2 ml) was added with a solution of Intermediate s-2 (362 mg) in methanol (3.2 ml) under ice cooling. This mixture was added dropwise with an aqueous solution (320  $\mu$ l) of sodium nitrite (84 mg, WAKO).

An aqueous solution (3.2 ml) of copper(II) bromide (270 mg, WAKO) was heated to 40°C, added dropwise with the previously obtained solution over 20 minutes, and stirred at the same temperature for 1.5 hours.

The reaction mixture was extracted with ethyl acetate (40 ml). The organic layer was washed successively with water and saturated brine, and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography (hexane:ethyl acetate = 9:1) to obtain the title compound (Intermediate s-3, 167 mg).

Synthesis of methyl 3-(3-bromo-4-fluorophenyl)acrylate (Intermediate s-4)

(Synthesis method SF)

According to the procedure described in the synthesis method of

Intermediate n-1 (Synthesis method SF) provided that the reaction was performed for 1 hour, 3-bromo-4-fluorocinnamic acid (3.30 g, LANC) and thionyl chloride (1.5 ml, WAKO) were reacted and treated to obtain the title compound (Intermediate n-25, 3.47 g).

Synthesis of methyl 3-[3-bromo-4-(4-methoxyphenylthio)phenyl]acrylate (Intermediate s-5) (Synthesis method SC)

A solution of Intermediate s-4 (259.1 mg) in DMSO (4 ml) was added with potassium carbonate (156.9 mg) and p-methoxythiophenol (148  $\mu$ l, TCI), and stirred at 70°C for 16 hours. The reaction mixture was extracted with ethyl acetate (30 ml), and then the organic layer was washed successively with water and saturated brine, and dried. Then, the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 8:1) to obtain the title compound (Intermediate s-5, 283.3 mg).

Synthesis of methyl 3-[3-bromo-4-(4-methoxyphenylthio)phenyl]propionate (Intermediate s-6) (Synthesis method SD2)

According to a procedure described in literature [D.J. Hart et al., Journal of Organic Chemistry (J. Org. Chem.), 1987, vol. 52, p.4665], a solution of Intermediate s-5 (579.1 mg) in dimethoxyethane (40 ml) was added with p-toluenesulfonhydrazide (1.99 g, TCI), and refluxed by heating at 110°C. Then, the reaction mixture was added dropwise with an aqueous solution (40 ml) of sodium acetate (1.54 g, WAKO) over 1 hour, and further stirred for 3 hours. The reaction mixture was extracted with dichloromethane (150 ml), and the organic layer was washed with water, and dried. Then, the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 7:1) to obtain the title compound (Intermediate s-6, 583.5 mg).

### Synthesis of 3-bromo-4-(cyclopentylthio)benzaldehyde (Intermediate s-23)

#### (Synthesis method SC)

A solution of 3-bromo-4-fluorobenzaldehyde (517.4 mg) in DMSO (8 ml) was added with potassium carbonate (514.9 mg) and cyclopentanethiol (250  $\mu$ l, TCI), and stirred at 90°C for 17 hours. The reaction mixture was extracted with ethyl acetate (50 ml), and the organic layer was washed successively with water and saturated brine, and dried. Then, the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 8:1) to obtain the title compound (Intermediate S-23, 644.7 mg).

### Synthesis of ethyl 3-[3-bromo-4-(cyclopentylthio)phenyl]acrylate (Intermediate s-24) (Synthesis method SE2)

A solution of Intermediate s-23 (243.7 mg) in 1,2-dimethoxyethane (8 ml) was added with ethyl diethylphosphonoacetate (300  $\mu$ l, TCI), and added with 60% sodium hydride (49.8 mg) under ice cooling. The reaction mixture was stirred for 10 minutes, then warmed to room temperature, and stirred for 1 hour. The reaction mixture was added with water (5 ml) for quenching, added with dichloromethane (30 ml) for extraction, and washed with saturated brine. The organic layer was dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 4:1) to obtain the title compound (Intermediate s-24, 286.2 mg).

Typical examples of the intermediates including those mentioned above that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification are shown in Table-Int. S-1 and Table-Int. S-2. In the tables, intermediate numbers are mentioned in the columns indicated as "Exp". In the tables, used methods among those



described above are mentioned in the columns of "Syn" with symbols, the starting compounds 1 are mentioned in the columns of "SM1", and the starting compounds 2 are mentioned in the columns of "SM2". Further, the compounds indicated as "Single" in the columns of "Single or Double" in Table-Int.S-1 are compounds in which two of the carbon atoms binding the benzene ring and carbonyl group in the compounds are bound with a single bond, and those indicated as "Double" in the same are compounds in which two of the carbon atoms binding the benzene ring and carbonyl group in the compounds are bound with a double bond.

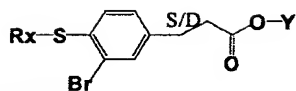


Table-Int.S-1

Exp.	Syn.	SM1	SM2	Rx-S	Y	Single or Double	LCMS		
							method	RTime	Mass
Ints-5	SC	Ints-4	4MeOPhSH	4MeOPhS	Me	Double	D	5.87	378(M <sup>+</sup> +1)
Ints-6	SD2	Ints-5		4MeOPhS	Me	Single	C		380(M <sup>+</sup> +1)
Ints-7	SC	Ints-4	2MeOPhSH	2MeOPhS	Me	Double	C		378(M <sup>+</sup> +1)
Ints-8	SD2	Ints-7		2MeOPhS	Me	Single	C		380(M <sup>+</sup> +1)
Ints-9	SC	Ints-4	3MeOPhSH	3MeOPhS	Me	Double	C		378(M <sup>+</sup> +1)
Ints-10	SD2	Ints-9		3MeOPhS	Me	Single	C		380(M <sup>+</sup> +1)
Ints-11	SC	Ints-4	2MePhSH	2MePhS	Me	Double	C		368(M <sup>+</sup> +1)
Ints-12	SD2	Ints-11		2MePhS	Me	Single	D	5.70	N.D
Ints-13	SC	Ints-4	3MePhSH	3MePhS	Me	Double	C		368(M <sup>+</sup> +1)
Ints-14	SD2	Ints-13		3MePhS	Me	Single	C		366(M <sup>+</sup> +1)
Ints-15	SC	Ints-4	4MePhSH	4MePhS	Me	Double	C		368(M <sup>+</sup> +1)
Ints-16	SD2	Ints-15		4MePhS	Me	Single	C		366(M <sup>+</sup> +1)
Ints-17	SC	Ints-4	2FPhSH	2FPhS	Me	Double	C		368(M <sup>+</sup> +1)
Ints-18	SD2	Ints-17		2FPhS	Me	Single	C		370(M <sup>+</sup> +1)
Ints-19	SC	Ints-4	3FPhSH	3FPhS	Me	Double	C		368(M <sup>+</sup> +1)
Ints-20	SD2	Ints-19		3FPhS	Me	Single	C		370(M <sup>+</sup> +1)
Ints-21	SC	Ints-4	4FPhSH	4FPhS	Me	Double	C		368(M <sup>+</sup> +1)
Ints-22	SD2	Ints-21		4FPhS	Me	Single	C		370(M <sup>+</sup> +1)
Ints-24	SE2	Ints-23		cPenS	Me	Double	D	6.35	340(M <sup>+</sup> +1)
Ints-25	SD2	Ints-24		cPenS	Me	Single	C		342(M <sup>+</sup> +1)
Ints-27	SE2	Ints-26		cHexS	Et	Double	C		354(M <sup>+</sup> +1)
Ints-28	SD2	Ints-27		cHexS	Et	Single	C		356(M <sup>+</sup> +1)
Ints-30	SE2	Ints-29		nPrS	Et	Double	C		328(M <sup>+</sup> +1)
Ints-31	SD2	Ints-30		nPrS	Et	Single	C		330(M <sup>+</sup> +1)
Ints-33	SE2	Ints-32		iPrS	Et	Double	C		328(M <sup>+</sup> +1)
Ints-34	SD2	Ints-33		iPrS	Et	Single	C		330(M <sup>+</sup> +1)
Ints-36	SE2	Ints-35		nBuS	Et	Double	C		328(M <sup>+</sup> +1)
Ints-37	SD2	Ints-36		nBuS	Et	Single	C		330(M <sup>+</sup> +1)
Ints-39	SE2	Ints-38		iBuS	Me	Double	D	5.86	330(M <sup>+</sup> +1)
Ints-40	SD2	Ints-39		iBuS	Me	Single	D	6.23	330(M <sup>+</sup> +1)
Ints-42	SE2	Ints-41		2PhEtS	Me	Double	D	6.18	376 (M <sup>+</sup> )
Ints-43	SD2	Ints-42		2PhEtS	Me	Single	D	6.21	378 (M <sup>+</sup> )
Ints-45	SE2	Ints-44		4MeOBnS	Et	Double	C		393 (M <sup>+</sup> +1)
Ints-46	SD2	Ints-45		4MeOBnS	Et	Single	C		395 (M <sup>+</sup> +1)
Ints-48	SE2	Ints-47		4FBnS	Et	Double	C		381 (M <sup>+</sup> +1)
Ints-49	SD2	Ints-48		4FBnS	Et	Single	C		383 (M <sup>+</sup> +1)
Ints-51	SE2	Ints-50		2MeBnS	Et	Double	C		377 (M <sup>+</sup> +1)
Ints-52	SD2	Ints-51		2MeBnS	Et	Single	C		379 (M <sup>+</sup> +1)

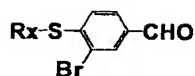


Table-Int.S-2

Exp.	Syn.	SM1	SM2	Rx-S	LCMS		
					method	RTime	Mass
Ints-23	SC		cPenSH	cPenS	C		286 (M <sup>+</sup> +1)
Ints-26	SC		cHexSH	cHexS	C		300 (M <sup>+</sup> +1)
Ints-29	SC		nPrSH	nPrS	C		260 (M <sup>+</sup> +1)
Ints-32	SC		iPrSH	iPrS	C		260 (M <sup>+</sup> +1)
Ints-35	SC		nBuSH	nBuS	C		274 (M <sup>+</sup> +1)
Ints-38	SC		iBuSH	iBuS	C		274 (M <sup>+</sup> +1)
Ints-41	SC		2PhEtSH	2PhEtS	C		322 (M <sup>+</sup> +1)
Ints-44	SC		4MeOBnSH	4MeOBnS	C		322 (M <sup>+</sup> +1)
Ints-47	SC		4FBnSH	4FBnS	C		326 (M <sup>+</sup> +1)
Ints-50	SC		2MeBnSH	2MeBnS	C		322 (M <sup>+</sup> +1)

[Example S-a-1]

Synthesis of methyl 3-[3-(naphthalen-2-yl)-4-(4-methylphenylthio)phenyl]propionate (Compound No. N-a-1) (Synthesis method SB)

A solution of Intermediate s-3 (146 mg) in toluene (2 ml) was added with 2-naphthaleneboronic acid (132.3 mg, TCI), 2 M aqueous sodium carbonate (600  $\mu$  ml), methanol (500  $\mu$  l), and tetrakis(triphenylphosphine) palladium(0) (henceforth abbreviated as "(Ph<sub>3</sub>P)<sub>4</sub>Pd", 38 mg, Nacalai Tesque), and stirred at 80°C for 14.5 hours. The reaction mixture was added with ethyl acetate (40 ml), and washed successively with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride and saturated brine. The organic layer was dried, and then the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography (hexane:ethyl acetate = 4:1) to obtain the title compound (Compound No. S-a-1, 78 mg).

[Example S-a-2]

Synthesis of 3-[3-(naphthalen-2-yl)-4-(4-methylphenylthio)phenyl]propionic acid (Compound No. S-a-2) (Synthesis method SA)

A solution of the compound of Example S-a-1 (51 mg) in methanol (5.0 ml) was added with 2 N aqueous sodium hydroxide (130  $\mu$  l), and stirred at 60°C for 2

hours. The reaction mixture was concentrated under reduced pressure, then made acidic with 5% aqueous hydrochloric acid under ice cooling, and then extracted with ethyl acetate (30 ml). The organic layer was washed with saturated brine, and dried, and then the solvent was evaporated under reduced pressure to obtain the title compound (Compound No. S-a-2, 47 mg).

[Example S-c-1]

Synthesis of methyl 3-[4-(4-methoxyphenylthio)-3-(naphthalen-2-yl)phenyl]propionate

(Compound No. S-c-1) (Synthesis method SD2)

According to the procedure described in the synthesis method of Intermediate s-6 (Synthesis method), the compound of Example S-b-1 (3.01 g), p-toluenesulfonhydrazide (430.1 mg), and sodium acetate (380.4 mg) were reacted and treated to obtain the title compound (Compound No. S-c-1, 95.1 mg).

[Examples S-a-1 to S-a-24, S-b-1 to S-b-138 and S-c-1 to S-c-138]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification including the examples described above are shown in Table-S-A-1, Table-S-B-1 to Table-S-B-3 and Table-S-C-1 to Table-S-C-3. In the tables, the compound numbers are mentioned in the columns indicated as "Exp.". In the tables, used methods among the aforementioned synthesis methods are shown in the columns of "Syn" with symbols, the starting compounds 1 are mentioned in the columns of "SM1", and the starting compounds 2 are mentioned in the columns of "SM2". The boronic acid reagents shown with the symbols of "BRA (number)" in the columns of "SM2" are those mentioned in Table-Ba-1 and Table-Ba-2.

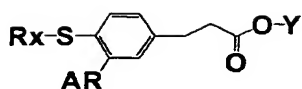


Table-S-A-1

Exp.	Syn	SM1	SM2	Rx	Y	AR	LCMS		
							method	RTime	Mass
S-a-1	SB	Ints-3	BRA1	4MePh	Me	2-Nap	C		413 (M <sup>+</sup> +1)
S-a-2	SA	S-a-1		4MePh	H	2-Nap	C		399 (M <sup>+</sup> +1)
S-a-3	SB	Ints-3	BRA2	4MePh	Me	5-Ind	C		402 (M <sup>+</sup> +1)
S-a-4	SA	S-a-3		4MePh	H	5-Ind	C		388 (M <sup>+</sup> +1)
S-a-5	SB	Ints-3	BRA3	4MePh	Me	1Me-5-Ind	C		416 (M <sup>+</sup> +1)
S-a-6	SA	S-a-5		4MePh	H	1Me-5-Ind	C		402 (M <sup>+</sup> +1)
S-a-7	SB	Ints-3	BRA4	4MePh	Me	1Et-5-Ind	C		430 (M <sup>+</sup> +1)
S-a-8	SA	S-a-7		4MePh	H	1Et-5-Ind	C		416 (M <sup>+</sup> +1)
S-a-9	SB	Ints-3	BRA5	4MePh	Me	5-1HIdz	C		403 (M <sup>+</sup> +1)
S-a-10	SA	S-a-9		4MePh	H	5-1HIdz	C		389 (M <sup>+</sup> +1)
S-a-11	SB	Ints-3	BRA6	4MePh	Me	1Me-5-1HIdz	C		417 (M <sup>+</sup> +1)
S-a-12	SA	S-a-11		4MePh	H	1Me-5-1HIdz	C		403 (M <sup>+</sup> +1)
S-a-13	SB	Ints-3	BRA7	4MePh	Me	1Et-5-1HIdz	C		431 (M <sup>+</sup> +1)
S-a-14	SA	S-a-13		4MePh	H	1Et-5-1HIdz	C		417 (M <sup>+</sup> +1)
S-a-15	SB	Ints-3	BRA8	4MePh	Me	2Me-5-2HIdz	C		417 (M <sup>+</sup> +1)
S-a-16	SA	S-a-15		4MePh	H	2Me-5-2HIdz	C		403 (M <sup>+</sup> +1)
S-a-17	SB	Ints-3	BRA9	4MePh	Me	5-Bzt	C		420 (M <sup>+</sup> +1)
S-a-18	SA	S-a-17		4MePh	H	5-Bzt	C		406 (M <sup>+</sup> +1)
S-a-19	SB	Ints-3	BRA10	4MePh	Me	3-Qu	C		414 (M <sup>+</sup> +1)
S-a-20	SA	S-a-19		4MePh	H	3-Qu	C		400 (M <sup>+</sup> +1)
S-a-21	SB	Ints-3	BRA11	4MePh	Me	6-Qu	C		414 (M <sup>+</sup> +1)
S-a-22	SA	S-a-21		4MePh	H	6-Qu	C		400 (M <sup>+</sup> +1)
S-a-23	SB	Ints-3	BRA12	4MePh	Me	6-IQ	C		414 (M <sup>+</sup> +1)
S-a-24	SA	N-a-23		4MePh	H	6-IQ	C		400 (M <sup>+</sup> +1)

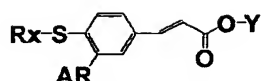


Table-S-B-1

Exp.	Syn	SM1	SM2	Rx	Y	AR	LCMS		
							method	RTime	Mass
S-b-1	SB	Ints-5	BRA1	4MeOPh	Me	2-Nap	C		427 (M <sup>+</sup> +1)
S-b-2	SA	S-b-1		4MeOPh	H	2-Nap	C		413 (M <sup>+</sup> +1)
S-b-3	SB	Ints-5	BRA2	4MeOPh	Me	5-Ind	C		416 (M <sup>+</sup> +1)
S-b-4	SA	S-b-3		4MeOPh	H	5-Ind	C		402 (M <sup>+</sup> +1)
S-b-5	SB	Ints-5	BRA3	4MeOPh	Me	1Me-5-Ind	C		430 (M <sup>+</sup> +1)
S-b-6	SA	S-b-5		4MeOPh	H	1Me-5-Ind	C		416 (M <sup>+</sup> +1)
S-b-7	SB	Ints-5	BRA4	4MeOPh	Me	1Et-5-Ind	C		444 (M <sup>+</sup> +1)
S-b-8	SA	S-b-7		4MeOPh	H	1Et-5-Ind	C		430 (M <sup>+</sup> +1)
S-b-9	SB	Ints-5	BRA5	4MeOPh	Me	5-1HIdz	C		417 (M <sup>+</sup> +1)
S-b-10	SA	S-b-9		4MeOPh	H	5-1HIdz	C		403 (M <sup>+</sup> +1)
S-b-11	SB	Ints-5	BRA6	4MeOPh	Me	1Me-5-1HIdz	C		431 (M <sup>+</sup> +1)
S-b-12	SA	S-b-11		4MeOPh	H	1Me-5-1HIdz	C		417 (M <sup>+</sup> +1)
S-b-13	SB	Ints-5	BRA7	4MeOPh	Me	1Et-5-1HIdz	C		445 (M <sup>+</sup> +1)
S-b-14	SA	S-b-13		4MeOPh	H	1Et-5-1HIdz	C		431 (M <sup>+</sup> +1)
S-b-15	SB	Ints-5	BRA8	4MeOPh	Me	2Me-5-2HIdz	C		431 (M <sup>+</sup> +1)
S-b-16	SA	S-b-15		4MeOPh	H	2Me-5-2HIdz	C		417 (M <sup>+</sup> +1)
S-b-17	SB	Ints-5	BRA9	4MeOPh	Me	5-Bzt	C		434 (M <sup>+</sup> +1)
S-b-18	SA	S-b-17		4MeOPh	H	5-Bzt	C		420 (M <sup>+</sup> +1)
S-b-19	SB	Ints-5	BRA10	4MeOPh	Me	3-Qu	C		428 (M <sup>+</sup> +1)
S-b-20	SA	S-b-19		4MeOPh	H	3-Qu	C		414 (M <sup>+</sup> +1)
S-b-21	SB	Ints-5	BRA11	4MeOPh	Me	6-Qu	C		428 (M <sup>+</sup> +1)
S-b-22	SA	S-b-21		4MeOPh	H	6-Qu	C		414 (M <sup>+</sup> +1)
S-b-23	SB	Ints-5	BRA12	4MeOPh	Me	6-IQ	C		428 (M <sup>+</sup> +1)
S-b-24	SA	S-b-23		4MeOPh	H	6-IQ	C		414 (M <sup>+</sup> +1)
S-b-25	SB	Ints-7	BRA1	2MeOPh	Me	2-Nap	C		427 (M <sup>+</sup> +1)
S-b-26	SA	S-b-25		2MeOPh	H	2-Nap	C		413 (M <sup>+</sup> +1)
S-b-27	SB	Ints-7	BRA2	2MeOPh	Me	5-Ind	C		416 (M <sup>+</sup> +1)
S-b-28	SA	S-b-27		2MeOPh	H	5-Ind	C		402 (M <sup>+</sup> +1)
S-b-29	SB	Ints-7	BRA5	2MeOPh	Me	5-1HIdz	C		417 (M <sup>+</sup> +1)
S-b-30	SA	S-b-29		2MeOPh	H	5-1HIdz	C		403 (M <sup>+</sup> +1)
S-b-31	SB	Ints-7	BRA10	2MeOPh	Me	3-Qu	C		428 (M <sup>+</sup> +1)
S-b-32	SA	S-b-31		2MeOPh	H	3-Qu	C		414 (M <sup>+</sup> +1)
S-b-33	SB	Ints-9	BRA1	3MeOPh	Me	2-Nap	C		427 (M <sup>+</sup> +1)
S-b-34	SA	S-b-33		3MeOPh	H	2-Nap	C		413 (M <sup>+</sup> +1)
S-b-35	SB	Ints-9	BRA3	3MeOPh	Me	1Me-5-Ind	C		430 (M <sup>+</sup> +1)
S-b-36	SA	S-b-35		3MeOPh	H	1Me-5-Ind	C		416 (M <sup>+</sup> +1)
S-b-37	SB	Ints-9	BRA6	3MeOPh	Me	1Me-5-1HIdz	C		431 (M <sup>+</sup> +1)
S-b-38	SA	S-b-37		3MeOPh	H	1Me-5-1HIdz	C		417 (M <sup>+</sup> +1)
S-b-39	SB	Ints-9	BRA11	3MeOPh	Me	6-Qu	C		428 (M <sup>+</sup> +1)
S-b-40	SA	S-b-39		3MeOPh	H	6-Qu	C		414 (M <sup>+</sup> +1)
S-b-41	SB	Ints-11	BRA2	2MePh	Me	5-Ind	C		400 (M <sup>+</sup> +1)
S-b-42	SA	S-b-41		2MePh	H	5-Ind	C		386 (M <sup>+</sup> +1)
S-b-43	SB	Ints-11	BRA3	2MePh	Me	1Me-5-Ind	C		414 (M <sup>+</sup> +1)
S-b-44	SA	S-b-43		2MePh	H	1Me-5-Ind	C		400 (M <sup>+</sup> +1)
S-b-45	SB	Ints-11	BRA5	2MePh	Me	5-1HIdz	C		401 (M <sup>+</sup> +1)
S-b-46	SA	S-b-45		2MePh	H	5-1HIdz	C		387 (M <sup>+</sup> +1)

Table-S-B-2

Exp.	Syn	SM1	SM2	Rx	Y	AR	LCMS		
							method	RTime	Mass
S-b-47	SB	Int.s-13	BRA3	3MePh	Me	1Me-5-Ind	C		414 (M <sup>+</sup> +1)
S-b-48	SA	N-b-47		3MePh	H	1Me-5-Ind	C		400 (M <sup>+</sup> +1)
S-b-49	SB	Int.s-13	BRA6	3MePh	Me	1Me-5-1HIdz	C		415 (M <sup>+</sup> +1)
S-b-50	SA	N-b-49		3MePh	H	1Me-5-1HIdz	C		401 (M <sup>+</sup> +1)
S-b-51	SB	Int.s-13	BRA9	3MePh	Me	5-Bzt	C		418 (M <sup>+</sup> +1)
S-b-52	SA	N-b-51		3MePh	H	5-Bzt	C		404 (M <sup>+</sup> +1)
S-b-53	SB	Int.s-15	BRA1	4MePh	Me	2-Nap	C		411 (M <sup>+</sup> +1)
S-b-54	SA	N-b-53		4MePh	H	2-Nap	C		397 (M <sup>+</sup> +1)
S-b-55	SB	Int.s-15	BRA2	4MePh	Me	5-Ind	C		400 (M <sup>+</sup> +1)
S-b-56	SA	N-b-55		4MePh	H	5-Ind	C		386 (M <sup>+</sup> +1)
S-b-57	SB	Int.s-15	BRA3	4MePh	Me	1Me-5-Ind	C		418 (M <sup>+</sup> +1)
S-b-58	SA	N-b-57		4MePh	H	1Me-5-Ind	C		404 (M <sup>+</sup> +1)
S-b-59	SB	Int.s-17	BRA5	2FPh	Me	5-1HIdz	C		404 (M <sup>+</sup> +1)
S-b-60	SA	N-b-59		2FPh	H	5-1HIdz	C		390 (M <sup>+</sup> +1)
S-b-61	SB	Int.s-17	BRA6	2FPh	Me	1Me-5-Ind	C		418 (M <sup>+</sup> +1)
S-b-62	SA	N-b-61		2FPh	H	1Me-5-Ind	C		404 (M <sup>+</sup> +1)
S-b-63	SB	Int.s-17	BRA11	2FPh	Me	6-Qu	C		415 (M <sup>+</sup> +1)
S-b-64	SA	N-b-63		2FPh	H	6-Qu	C		401 (M <sup>+</sup> +1)
S-b-65	SB	Int.s-19	BRA1	3FPh	Me	2-Nap	C		415 (M <sup>+</sup> +1)
S-b-66	SA	N-b-65		3FPh	H	2-Nap	C		401 (M <sup>+</sup> +1)
S-b-67	SB	Int.s-19	BRA2	3FPh	Me	5-Ind	C		403 (M <sup>+</sup> +1)
S-b-68	SA	N-b-67		3FPh	H	5-Ind	C		389 (M <sup>+</sup> +1)
S-b-69	SB	Int.s-19	BRA6	3FPh	Me	1Me-5-1HIdz	C		418 (M <sup>+</sup> +1)
S-b-70	SA	N-b-69		3FPh	H	1Me-5-1HIdz	C		404 (M <sup>+</sup> +1)
S-b-71	SB	Int.s-21	BRA3	4FPh	Me	1Me-5-Ind	C		418 (M <sup>+</sup> +1)
S-b-72	SA	N-b-71		4FPh	H	1Me-5-Ind	C		404 (M <sup>+</sup> +1)
S-b-73	SB	Int.s-21	BRA5	4FPh	Me	5-1HIdz	C		404 (M <sup>+</sup> +1)
S-b-74	SA	N-b-73		4FPh	H	5-1HIdz	C		390 (M <sup>+</sup> +1)
S-b-75	SB	Int.s-21	BRA10	4FPh	Me	3-Qu	C		415 (M <sup>+</sup> +1)
S-b-76	SA	N-b-75		4FPh	H	3-Qu	C		401 (M <sup>+</sup> +1)
S-b-77	SB	Int.s-24	BRA1	cPen	Me	2-Nap	C		389 (M <sup>+</sup> +1)
S-b-78	SA	N-b-77		cPen	H	2-Nap	C		375 (M <sup>+</sup> +1)
S-b-79	SB	Int.s-24	BRA2	cPen	Me	5-Ind	C		378 (M <sup>+</sup> +1)
S-b-80	SA	N-b-79		cPen	H	5-Ind	C		364 (M <sup>+</sup> +1)
S-b-81	SB	Int.s-24	BRA6	cPen	Me	1Me-5-1HIdz	C		407 (M <sup>+</sup> +1)
S-b-82	SA	N-b-81		cPen	H	1Me-5-1HIdz	C		393 (M <sup>+</sup> +1)
S-b-83	SB	Int.s-27	BRA3	cHex	Et	1Me-5-Ind	C		406 (M <sup>+</sup> +1)
S-b-84	SA	N-b-83		cHex	H	1Me-5-Ind	C		392 (M <sup>+</sup> +1)
S-b-85	SB	Int.s-27	BRA5	cHex	Et	5-1HIdz	C		393 (M <sup>+</sup> +1)
S-b-86	SA	N-b-85		cHex	H	5-1HIdz	C		379 (M <sup>+</sup> +1)
S-b-87	SB	Int.s-27	BRA12	cHex	Et	6-Qu	C		363 (M <sup>+</sup> +1)
S-b-88	SA	N-b-87		cHex	H	6-Qu	C		349 (M <sup>+</sup> +1)
S-b-89	SB	Int.s-30	BRA1	nPr	Et	2-Nap	C		362 (M <sup>+</sup> +1)
S-b-90	SA	N-b-89		nPr	H	2-Nap	C		348 (M <sup>+</sup> +1)
S-b-91	SB	Int.s-30	BRA2	nPr	Et	5-Ind	C		351 (M <sup>+</sup> +1)
S-b-92	SA	N-b-91		nPr	H	5-Ind	C		337 (M <sup>+</sup> +1)

Table-S-B-3

Exp.	Syn	SM1	SM2	Rx	Y	AR	LCMS		
							method	RTime	Mass
S-b-93	SB	Int.s-30	BRA6	nPr	Et	1Me-5-1HIdz	C		366 (M <sup>+</sup> +1)
S-b-94	SA	N-b-93		nPr	H	1Me-5-1HIdz	C		352 (M <sup>+</sup> +1)
S-b-95	SB	Int.s-33	BRA1	iPr	Et	2-Nap	C		362 (M <sup>+</sup> +1)
S-b-96	SA	N-b-95		iPr	H	2-Nap	C		348 (M <sup>+</sup> +1)
S-b-97	SB	Int.s-33	BRA3	iPr	Et	1Me-5-Ind	C		365 (M <sup>+</sup> +1)
S-b-98	SA	N-b-97		iPr	H	1Me-5-Ind	C		351 (M <sup>+</sup> +1)
S-b-99	SB	Int.s-33	BRA5	iPr	Et	5-1HIdz	C		352 (M <sup>+</sup> +1)
S-b-100	SA	N-b-99		iPr	H	5-1HIdz	C		338 (M <sup>+</sup> +1)
S-b-101	SB	Int.s-36	BRA2	nBu	Et	5-Ind	C		366 (M <sup>+</sup> +1)
S-b-102	SA	N-b-101		nBu	H	5-Ind	C		352 (M <sup>+</sup> +1)
S-b-103	SB	Int.s-36	BRA5	nBu	Et	5-1HIdz	C		366 (M <sup>+</sup> +1)
S-b-104	SA	N-b-103		nBu	H	5-1HIdz	C		352 (M <sup>+</sup> +1)
S-b-105	SB	Int.s-36	BRA11	nBu	Et	6-Qu	C		378 (M <sup>+</sup> +1)
S-b-106	SA	N-b-105		nBu	H	6-Qu	C		364 (M <sup>+</sup> +1)
S-b-107	SB	Int.s-39	BRA1	iBu	Me	2-Nap	C		377 (M <sup>+</sup> +1)
S-b-108	SA	N-b-107		iBu	H	2-Nap	C		363 (M <sup>+</sup> +1)
S-b-109	SB	Int.s-39	BRA3	iBu	Me	1Me-5-Ind	C		380 (M <sup>+</sup> +1)
S-b-110	SA	N-b-109		iBu	H	1Me-5-Ind	C		366 (M <sup>+</sup> +1)
S-b-111	SB	Int.s-39	BRA5	iBu	Me	5-1HIdz	C		366 (M <sup>+</sup> +1)
S-b-112	SA	N-b-111		iBu	H	5-1HIdz	C		352 (M <sup>+</sup> +1)
S-b-113	SB	Int.s-39	BRA6	iBu	Me	1Me-5-1HIdz	C		381 (M <sup>+</sup> +1)
S-b-114	SA	N-b-113		iBu	H	1Me-5-1HIdz	C		367 (M <sup>+</sup> +1)
S-b-115	SB	Int.s-42	BRA1	PhEt	Me	2-Nap	C		425 (M <sup>+</sup> +1)
S-b-116	SA	N-b-115		PhEt	H	2-Nap	C		411 (M <sup>+</sup> +1)
S-b-117	SB	Int.s-42	BRA2	PhEt	Me	5-Ind	C		414 (M <sup>+</sup> +1)
S-b-118	SA	N-b-117		PhEt	H	5-Ind	C		400 (M <sup>+</sup> +1)
S-b-119	SB	Int.s-42	BRA3	PhEt	Me	1Me-5-Ind	C		428 (M <sup>+</sup> +1)
S-b-120	SA	N-b-119		PhEt	H	1Me-5-Ind	C		414 (M <sup>+</sup> +1)
S-b-121	SB	Int.s-45	BRA1	4MeOBn	Et	2-Nap	C		441 (M <sup>+</sup> +1)
S-b-122	SA	N-b-121		4MeOBn	H	2-Nap	C		427 (M <sup>+</sup> +1)
S-b-123	SB	Int.s-45	BRA5	4MeOBn	Et	5-1HIdz	C		431 (M <sup>+</sup> +1)
S-b-124	SA	N-b-123		4MeOBn	H	5-1HIdz	C		417 (M <sup>+</sup> +1)
S-b-125	SB	Int.s-45	BRA6	4MeOBn	Et	1Me-5-1HIdz	C		431 (M <sup>+</sup> +1)
S-b-126	SA	N-b-125		4MeOBn	H	1Me-5-1HIdz	C		417 (M <sup>+</sup> +1)
S-b-127	SB	Int.s-48	BRA1	4FBn	Et	2-Nap	C		429 (M <sup>+</sup> +1)
S-b-128	SA	N-b-127		4FBn	H	2-Nap	C		415 (M <sup>+</sup> +1)
S-b-129	SB	Int.s-48	BRA2	4FBn	Et	5-Ind	C		418 (M <sup>+</sup> +1)
S-b-130	SA	N-b-129		4FBn	H	5-Ind	C		404 (M <sup>+</sup> +1)
S-b-131	SB	Int.s-48	BRA6	4FBn	Et	1Me-5-1HIdz	C		418 (M <sup>+</sup> +1)
S-b-132	SA	N-b-131		4FBn	H	1Me-5-1HIdz	C		404 (M <sup>+</sup> +1)
S-b-133	SB	Int.s-51	BRA3	2MeBn	Et	1Me-5-Ind	C		428 (M <sup>+</sup> +1)
S-b-134	SA	N-b-133		2MeBn	H	1Me-5-Ind	C		414 (M <sup>+</sup> +1)
S-b-135	SB	Int.s-51	BRA5	2MeBn	Et	5-1HIdz	C		415 (M <sup>+</sup> +1)
S-b-136	SA	N-b-135		2MeBn	H	5-1HIdz	C		401 (M <sup>+</sup> +1)
S-b-137	SB	Int.s-51	BRA10	2MeBn	Et	3-Qu	C		426 (M <sup>+</sup> +1)
S-b-138	SA	N-b-137		2MeBn	H	3-Qu	C		412 (M <sup>+</sup> +1)



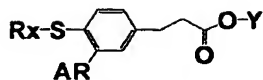


Table-S-C-1

Exp.	Syn	SM1	SM2	Rx	Y	AR	LCMS		
							method	RTime	Mass
S-c-1	SB	Ints-6	BRA1	4MeOPh	Me	2-Nap	C		429 (M <sup>+</sup> +1)
S-c-2	SA	S-c-1		4MeOPh	H	2-Nap	C		415 (M <sup>+</sup> +1)
S-c-3	SB	Ints-6	BRA2	4MeOPh	Me	5-Ind	C		418 (M <sup>+</sup> +1)
S-c-4	SA	S-c-3		4MeOPh	H	5-Ind	C		404 (M <sup>+</sup> +1)
S-c-5	SB	Ints-6	BRA3	4MeOPh	Me	1Me-5-Ind	C		432 (M <sup>+</sup> +1)
S-c-6	SA	S-c-5		4MeOPh	H	1Me-5-Ind	C		418 (M <sup>+</sup> +1)
S-c-7	SB	Ints-6	BRA4	4MeOPh	Me	1Et-5-Ind	C		446 (M <sup>+</sup> +1)
S-c-8	SA	S-c-7		4MeOPh	H	1Et-5-Ind	C		432 (M <sup>+</sup> +1)
S-c-9	SB	Ints-6	BRA5	4MeOPh	Me	5-1HIdz	C		419 (M <sup>+</sup> +1)
S-c-10	SA	S-c-9		4MeOPh	H	5-1HIdz	C		405 (M <sup>+</sup> +1)
S-c-11	SB	Ints-6	BRA6	4MeOPh	Me	1Me-5-1HIdz	C		433 (M <sup>+</sup> +1)
S-c-12	SA	S-c-11		4MeOPh	H	1Me-5-1HIdz	C		419 (M <sup>+</sup> +1)
S-c-13	SB	Ints-6	BRA7	4MeOPh	Me	1Et-5-1HIdz	C		447 (M <sup>+</sup> +1)
S-c-14	SA	S-c-13		4MeOPh	H	1Et-5-1HIdz	C		433 (M <sup>+</sup> +1)
S-c-15	SB	Ints-6	BRA8	4MeOPh	Me	2Me-5-2HIdz	C		433 (M <sup>+</sup> +1)
S-c-16	SA	S-c-15		4MeOPh	H	2Me-5-2HIdz	C		419 (M <sup>+</sup> +1)
S-c-17	SB	Ints-6	BRA9	4MeOPh	Me	5-Bzt	C		436 (M <sup>+</sup> +1)
S-c-18	SA	S-c-17		4MeOPh	H	5-Bzt	C		422 (M <sup>+</sup> +1)
S-c-19	SB	Ints-6	BRA10	4MeOPh	Me	3-Qu	C		430 (M <sup>+</sup> +1)
S-c-20	SA	S-c-19		4MeOPh	H	3-Qu	C		416 (M <sup>+</sup> +1)
S-c-21	SB	Ints-6	BRA11	4MeOPh	Me	6-Qu	C		430 (M <sup>+</sup> +1)
S-c-22	SA	S-c-21		4MeOPh	H	6-Qu	C		416 (M <sup>+</sup> +1)
S-c-23	SB	Ints-6	BRA12	4MeOPh	Me	6-IQ	C		430 (M <sup>+</sup> +1)
S-c-24	SA	S-c-23		4MeOPh	H	6-IQ	C		416 (M <sup>+</sup> +1)
S-c-25	SB	Ints-8	BRA1	2MeOPh	Me	2-Nap	C		429 (M <sup>+</sup> +1)
S-c-26	SA	S-c-25		2MeOPh	H	2-Nap	C		415 (M <sup>+</sup> +1)
S-c-27	SB	Ints-8	BRA3	2MeOPh	Me	1Me-5-Ind	C		431 (M <sup>+</sup> +1)
S-c-28	SA	S-c-27		2MeOPh	H	1Me-5-Ind	C		418 (M <sup>+</sup> +1)
S-c-29	SB	Ints-8	BRA5	2MeOPh	Me	5-1HIdz	C		419 (M <sup>+</sup> +1)
S-c-30	SA	S-c-29		2MeOPh	H	5-1HIdz	C		405 (M <sup>+</sup> +1)
S-c-31	SB	Ints-8	BRA10	2MeOPh	Me	3-Qu	C		430 (M <sup>+</sup> +1)
S-c-32	SA	S-c-31		2MeOPh	H	3-Qu	C		416 (M <sup>+</sup> +1)
S-c-33	SB	Ints-10	BRA1	3MeOPh	Me	2-Nap	C		429 (M <sup>+</sup> +1)
S-c-34	SA	S-c-33		3MeOPh	H	2-Nap	C		415 (M <sup>+</sup> +1)
S-c-35	SB	Ints-10	BRA2	3MeOPh	Me	5-Ind	C		418 (M <sup>+</sup> +1)
S-c-36	SA	S-c-35		3MeOPh	H	5-Ind	C		403 (M <sup>+</sup> +1)
S-c-37	SB	Ints-10	BRA6	3MeOPh	Me	1Me-5-1HIdz	C		433 (M <sup>+</sup> +1)
S-c-38	SA	S-c-37		3MeOPh	H	1Me-5-1HIdz	C		419 (M <sup>+</sup> +1)
S-c-39	SB	Ints-10	BRA11	3MeOPh	Me	6-Qu	C		430 (M <sup>+</sup> +1)
S-c-40	SA	S-c-39		3MeOPh	H	6-Qu	C		416 (M <sup>+</sup> +1)
S-c-41	SB	Ints-12	BRA3	2MePh	Me	1-Me-5-Ind	C		402 (M <sup>+</sup> +1)
S-c-42	SA	S-c-41		2MePh	H	1-Me-5-Ind	C		388 (M <sup>+</sup> +1)
S-c-43	SB	Ints-12	BRA5	2MePh	Me	5-1HIdz	C		416 (M <sup>+</sup> +1)
S-c-44	SA	S-c-43		2MePh	H	5-1HIdz	C		402 (M <sup>+</sup> +1)
S-c-45	SB	Ints-12	BRA6	2MePh	Me	1-Me-5-1HIdz	C		417 (M <sup>+</sup> +1)
S-c-46	SA	S-c-45		2MePh	H	1-Me-5-1HIdz	C		403 (M <sup>+</sup> +1)

Table-S-C-2

Exp.	Syn	SM1	SM2	Rx	Y	AR	LCMS		
							method	RTime	Mass
S-c-47	SB	Ints-14	BRA3	3MePh	Me	1-Me-5-Ind	C		416 (M <sup>+</sup> +1)
S-c-48	SA	N-c-47		3MePh	H	1-Me-5-Ind	C		402 (M <sup>+</sup> +1)
S-c-49	SB	Ints-14	BRA6	3MePh	Me	1-Me-5-1HIdz	C		417 (M <sup>+</sup> +1)
S-c-50	SA	N-c-49		3MePh	H	1-Me-5-1HIdz	C		403 (M <sup>+</sup> +1)
S-c-51	SB	Ints-14	BRA9	3MePh	Me	5-Bzt	C		420 (M <sup>+</sup> +1)
S-c-52	SA	N-c-51		3MePh	H	5-Bzt	C		406 (M <sup>+</sup> +1)
S-c-53	SB	Ints-16	BRA1	4MePh	Me	2-Nap	C		413 (M <sup>+</sup> +1)
S-c-54	SA	N-c-53		4MePh	H	2-Nap	C		399 (M <sup>+</sup> +1)
S-c-55	SB	Ints-16	BRA2	4MePh	Me	5-Ind	C		402 (M <sup>+</sup> +1)
S-c-56	SA	N-c-55		4MePh	H	5-Ind	C		388 (M <sup>+</sup> +1)
S-c-57	SB	Ints-16	BRA3	4MePh	Me	1Me-5-Ind	C		420 (M <sup>+</sup> +1)
S-c-58	SA	N-c-57		4MePh	H	1Me-5-Ind	C		406 (M <sup>+</sup> +1)
S-c-59	SB	Ints-18	BRA5	2FPh	Me	5-1HIdz	C		406 (M <sup>+</sup> +1)
S-c-60	SA	N-c-59		2FPh	H	5-1HIdz	C		392 (M <sup>+</sup> +1)
S-c-61	SB	Ints-18	BRA6	2FPh	Me	1Me-5-1HInd	C		420 (M <sup>+</sup> +1)
S-c-62	SA	N-c-61		2FPh	H	1Me-5-1HInd	C		406 (M <sup>+</sup> +1)
S-c-63	SB	Ints-18	BRA11	2FPh	Me	6-Qu	C		418 (M <sup>+</sup> +1)
S-c-64	SA	N-c-63		2FPh	H	6-Qu	C		404 (M <sup>+</sup> +1)
S-c-65	SB	Ints-20	BRA1	3FPh	Me	2-Nap	C		417 (M <sup>+</sup> +1)
S-c-66	SA	N-c-65		3FPh	H	2-Nap	C		403 (M <sup>+</sup> +1)
S-c-67	SB	Ints-20	BRA2	3FPh	Me	5-Ind	C		405 (M <sup>+</sup> +1)
S-c-68	SA	N-c-67		3FPh	H	5-Ind	C		391 (M <sup>+</sup> +1)
S-c-69	SB	Ints-20	BRA6	3FPh	Me	1Me-5-1HIdz	C		421 (M <sup>+</sup> +1)
S-c-70	SA	N-c-69		3FPh	H	1Me-5-1HIdz	C		407 (M <sup>+</sup> +1)
S-c-71	SB	Ints-22	BRA3	4FPh	Me	1Me-5-Ind	C		420 (M <sup>+</sup> +1)
S-c-72	SA	N-c-71		4FPh	H	1Me-5-Ind	C		406 (M <sup>+</sup> +1)
S-c-73	SB	Ints-22	BRA5	4FPh	Me	5-1HIdz	C		406 (M <sup>+</sup> +1)
S-c-74	SA	N-c-73		4FPh	H	5-1HIdz	C		392 (M <sup>+</sup> +1)
S-c-75	SB	Ints-22	BRA10	4FPh	Me	3-Qu	C		418 (M <sup>+</sup> +1)
S-c-76	SA	N-c-75		4FPh	H	3-Qu	C		404 (M <sup>+</sup> +1)
S-c-77	SB	Ints-25	BRA1	cPen	Et	2-Nap	C		391 (M <sup>+</sup> +1)
S-c-78	SA	N-c-77		cPen	H	2-Nap	C		377 (M <sup>+</sup> +1)
S-c-79	SB	Ints-25	BRA2	cPen	Et	5-Ind	C		380 (M <sup>+</sup> +1)
S-c-80	SA	N-c-79		cPen	H	5-Ind	C		366 (M <sup>+</sup> +1)
S-c-81	SB	Ints-25	BRA6	cPen	Et	1Me-5-1HIdz	C		409 (M <sup>+</sup> +1)
S-c-82	SA	N-c-81		cPen	H	1Me-5-1HIdz	C		395 (M <sup>+</sup> +1)
S-c-83	SB	Ints-28	BRA3	cHex	Et	1Me-5-Ind	C		408 (M <sup>+</sup> +1)
S-c-84	SA	N-c-83		cHex	H	1Me-5-Ind	C		394 (M <sup>+</sup> +1)
S-c-85	SB	Ints-28	BRA5	cHex	Et	5-1HIdz	C		395 (M <sup>+</sup> +1)
S-c-86	SA	N-c-85		cHex	H	5-1HIdz	C		381 (M <sup>+</sup> +1)
S-c-87	SB	Ints-28	BRA12	cHex	Et	6-Qu	C		365 (M <sup>+</sup> +1)
S-c-88	SA	N-c-87		cHex	H	6-Qu	C		351 (M <sup>+</sup> +1)
S-c-89	SB	Ints-31	BRA1	nPr	Et	2-Nap	C		365 (M <sup>+</sup> +1)
S-c-90	SA	N-c-89		nPr	H	2-Nap	C		351 (M <sup>+</sup> +1)
S-c-91	SB	Ints-31	BRA2	nPr	Et	5-Ind	C		353 (M <sup>+</sup> +1)
S-c-92	SA	N-c-91		nPr	H	5-Ind	C		339 (M <sup>+</sup> +1)

Table-S-C-3

Exp.	Syn	SM1	SM2	Rx	Y	AR	LCMS		
							method	RTime	Mass
S-c-93	SB	Ints-31	BRA6	nPr	Et	1Me-5-1HIdz	C		368 (M <sup>+</sup> +1)
S-c-94	SA	N-c-93		nPr	H	1Me-5-1HIdz	C		354 (M <sup>+</sup> +1)
S-c-95	SB	Ints-34	BRA3	iPr	Et	1Me-5-Ind	C		368 (M <sup>+</sup> +1)
S-c-96	SA	N-c-95		iPr	H	1Me-5-Ind	C		354 (M <sup>+</sup> +1)
S-c-97	SB	Ints-34	BRA5	iPr	Et	5-1HIdz	C		354 (M <sup>+</sup> +1)
S-c-98	SA	N-c-97		iPr	H	5-1HIdz	C		340 (M <sup>+</sup> +1)
S-c-99	SB	Ints-34	BRA12	iPr	Et	6-IQ	C		380 (M <sup>+</sup> +1)
S-c-100	SA	N-c-99		iPr	H	6-IQ	C		366 (M <sup>+</sup> +1)
S-c-101	SB	Ints-37	BRA1	nBu	Et	2-Nap	C		379 (M <sup>+</sup> +1)
S-c-102	SA	N-c-101		nBu	H	2-Nap	C		365 (M <sup>+</sup> +1)
S-c-103	SB	Ints-37	BRA2	nBu	Et	5-Ind	C		368 (M <sup>+</sup> +1)
S-c-104	SA	N-c-103		nBu	H	5-Ind	C		354 (M <sup>+</sup> +1)
S-c-105	SB	Ints-37	BRA6	nBu	Et	1Me-5-1HIdz	C		383 (M <sup>+</sup> +1)
S-c-106	SA	N-c-105		nBu	H	1Me-5-1HIdz	C		369 (M <sup>+</sup> +1)
S-c-107	SB	Ints-40	BRA1	iBu	Et	2-Nap	C		379 (M <sup>+</sup> +1)
S-c-108	SA	N-c-107		iBu	H	2-Nap	C		365 (M <sup>+</sup> +1)
S-c-109	SB	Ints-40	BRA3	iBu	Et	1Me-5-Ind	C		382 (M <sup>+</sup> +1)
S-c-110	SA	N-c-109		iBu	H	1Me-5-Ind	C		368 (M <sup>+</sup> +1)
S-c-111	SB	Ints-40	BRA5	iBu	Et	5-1HIdz	C		369 (M <sup>+</sup> +1)
S-c-112	SA	N-c-111		iBu	H	5-1HIdz	C		355 (M <sup>+</sup> +1)
S-c-113	SB	Ints-40	BRA6	iBu	Et	1Me-5-1HIdz	C		383 (M <sup>+</sup> +1)
S-c-114	SA	N-c-113		iBu	H	1Me-5-1HIdz	C		369 (M <sup>+</sup> +1)
S-c-115	SB	Ints-43	BRA1	PhEt	Et	2-Nap	C		427 (M <sup>+</sup> +1)
S-c-116	SA	N-c-115		PhEt	H	2-Nap	C		413 (M <sup>+</sup> +1)
S-c-117	SB	Ints-43	BRA2	PhEt	Et	5-Ind	C		416 (M <sup>+</sup> +1)
S-c-118	SA	N-c-117		PhEt	H	5-Ind	C		402 (M <sup>+</sup> +1)
S-c-119	SB	Ints-43	BRA6	PhEt	Et	1Me-5-1HIdz	C		431 (M <sup>+</sup> +1)
S-c-120	SA	N-c-119		PhEt	H	1Me-5-1HIdz	C		417 (M <sup>+</sup> +1)
S-c-121	SB	Ints-46	BRA1	4MeOBn	Et	2-Nap	C		443 (M <sup>+</sup> +1)
S-c-122	SA	N-c-121		4MeOBn	H	2-Nap	C		429 (M <sup>+</sup> +1)
S-c-123	SB	Ints-46	BRA3	4MeOBn	Et	1Me-5-Ind	C		446 (M <sup>+</sup> +1)
S-c-124	SA	N-c-123		4MeOBn	H	1Me-5-Ind	C		432 (M <sup>+</sup> +1)
S-c-125	SB	Ints-46	BRA5	4MeOBn	Et	5-1HIdz	C		419 (M <sup>+</sup> +1)
S-c-126	SA	N-c-125		4MeOBn	H	5-1HIdz	C		405 (M <sup>+</sup> +1)
S-c-127	SB	Ints-49	BRA1	4FBn	Et	2-Nap	C		431 (M <sup>+</sup> +1)
S-c-128	SA	N-c-127		4FBn	H	2-Nap	C		417 (M <sup>+</sup> +1)
S-c-129	SB	Ints-49	BRA2	4FBn	Et	5-Ind	C		420 (M <sup>+</sup> +1)
S-c-130	SA	N-c-129		4FBn	H	5-Ind	C		406 (M <sup>+</sup> +1)
S-c-131	SB	Ints-49	BRA5	4FBn	Et	5-1HIdz	C		406 (M <sup>+</sup> +1)
S-c-132	SA	N-c-131		4FBn	H	5-1HIdz	C		392 (M <sup>+</sup> +1)
S-c-133	SB	Ints-52	BRA3	2MeBn	Et	1Me-5-Ind	C		430 (M <sup>+</sup> +1)
S-c-134	SA	N-c-133		2MeBn	H	1Me-5-Ind	C		416 (M <sup>+</sup> +1)
S-c-135	SB	Ints-52	BRA6	2MeBn	Et	1Me-5-1HIdz	C		431 (M <sup>+</sup> +1)
S-c-136	SA	N-c-135		2MeBn	H	1Me-5-1HIdz	C		417 (M <sup>+</sup> +1)
S-c-137	SB	Ints-52	BRA11	2MeBn	Et	6-Qu	C		428 (M <sup>+</sup> +1)
S-c-138	SA	N-c-137		2MeBn	H	6-Qu	C		414 (M <sup>+</sup> +1)

[Example S-d-1]

Synthesis of ethyl 3-[4-[(4-methoxyphenyl)methylsulfinyl]-3-(naphthalen-2-

yl)phenyl}propionate (Compound No. S-d-1) (Synthesis method SG)

A solution of the compound of Example S-c-121 (130.9 mg) in dichloromethane (4 ml) was added with 3-chloroperoxybenzoic acid (60.0 mg, TCI), and stirred at room temperature for 1.5 hours. The reaction mixture was added with water (10 ml), extracted with dichloromethane (20 ml), and then washed with saturated brine. The organic layer was dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, chloroform:methanol = 30:1) to obtain the title compound (Compound No. S-d-1, 108.7 mg).

[Example S-d-7]

Synthesis of ethyl 3-{4-[(4-methoxyphenyl)methylsulfonyl]-3-(naphthalen-2-yl)phenyl}propionate (Compound No. S-d-7) (Synthesis method SG)

A solution of the compound of Example S-c-121 (53.1 mg) in dichloromethane (3 ml) was added with 3-chloroperoxybenzoic acid (74.5 mg, TCI), and stirred at room temperature for 5 hours. The reaction mixture was added with water (10 ml), extracted with dichloromethane (20 ml), and then washed with saturated brine. The organic layer was dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 4:1) to obtain the title compound (Compound No. S-d-7, 48.1 mg).

[Examples S-d-1 to S-d-36]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification including the examples described above are shown in Table-S-D-1. In the tables, the compound numbers are mentioned in the columns indicated as "Exp.". In the tables, used methods among the aforementioned synthesis methods are shown in the columns of "Syn" with

symbols, and the starting compounds 1 are mentioned in the columns of "SM1".

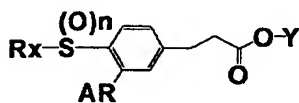


Table-S-D-1

Exp.	Syn	SM1	RS(O) <sub>n</sub>	Y	AR	LCMS		
						method	RTime	Mass
S-d-1	SG	S-c-121	4MeOBnSO	Et	2-Nap	C		473 (M <sup>+</sup> +1)
S-d-2	SA	S-d-1	4MeOBnSO	H	2-Nap	C		445 (M <sup>+</sup> +1)
S-d-3	SG	S-c-123	4MeOBnSO	Et	1Me-5-Ind	C		476 (M <sup>+</sup> +1)
S-d-4	SA	S-d-3	4MeOBnSO	H	1Me-5-Ind	C		448 (M <sup>+</sup> +1)
S-d-5	SG	S-c-125	4MeOBnSO	Et	5-1HIdz	C		463 (M <sup>+</sup> +1)
S-d-6	SA	S-d-5	4MeOBnSO	H	5-1HIdz	C		435 (M <sup>+</sup> +1)
S-d-7	SG	S-c-121	4MeOBnSO <sub>2</sub>	Et	2-Nap	C		489 (M <sup>+</sup> +1)
S-d-8	SA	S-d-7	4MeOBnSO <sub>2</sub>	H	2-Nap	C		461 (M <sup>+</sup> +1)
S-d-9	SG	S-c-123	4MeOBnSO <sub>2</sub>	Et	1Me-5-Ind	C		492 (M <sup>+</sup> +1)
S-d-10	SA	S-d-9	4MeOBnSO <sub>2</sub>	H	1Me-5-Ind	C		464 (M <sup>+</sup> +1)
S-d-11	SG	S-c-125	4MeOBnSO <sub>2</sub>	Et	5-1HIdz	C		479 (M <sup>+</sup> +1)
S-d-12	SA	S-d-11	4MeOBnSO <sub>2</sub>	H	5-1HIdz	C		451 (M <sup>+</sup> +1)
S-d-13	SG	S-c-77	cPenSO	Et	2-Nap	C		421 (M <sup>+</sup> +1)
S-d-14	SA	S-d-13	cPenSO	H	2-Nap	C		393 (M <sup>+</sup> +1)
S-d-15	SG	S-c-79	cPenSO	Et	5-Ind	C		410 (M <sup>+</sup> +1)
S-d-16	SA	S-d-15	cPenSO	H	5-Ind	C		381 (M <sup>+</sup> +1)
S-d-17	SG	S-c-81	cPenSO	Et	1Me-5-1HIdz	C		425 (M <sup>+</sup> +1)
S-d-18	SA	S-d-17	cPenSO	H	1Me-5-1HIdz	C		397 (M <sup>+</sup> +1)
S-d-19	SG	S-c-77	cPenSO <sub>2</sub>	Et	2-Nap	C		437 (M <sup>+</sup> +1)
S-d-20	SA	S-d-19	cPenSO <sub>2</sub>	H	2-Nap	C		409 (M <sup>+</sup> +1)
S-d-21	SG	S-c-79	cPenSO <sub>2</sub>	Et	5-Ind	C		426 (M <sup>+</sup> +1)
S-d-22	SA	S-d-21	cPenSO <sub>2</sub>	H	5-Ind	C		397 (M <sup>+</sup> +1)
S-d-23	SG	S-c-81	cPenSO <sub>2</sub>	Et	1Me-5-1HIdz	C		441 (M <sup>+</sup> +1)
S-d-24	SA	S-d-23	cPenSO <sub>2</sub>	H	1Me-5-1HIdz	C		413 (M <sup>+</sup> +1)
S-d-25	SG	S-c-101	nBuSO	Et	2-Nap	C		409 (M <sup>+</sup> +1)
S-d-26	SA	S-d-25	nBuSO	H	2-Nap	C		377 (M <sup>+</sup> +1)
S-d-27	SG	S-c-103	nBuSO	Et	5-Ind	C		398 (M <sup>+</sup> +1)
S-d-28	SA	S-d-27	nBuSO	H	5-Ind	C		365 (M <sup>+</sup> +1)
S-d-29	SG	S-c-105	nBuSO	Et	1Me-5-1HIdz	C		413 (M <sup>+</sup> +1)
S-d-30	SA	S-d-29	nBuSO	H	1Me-5-1HIdz	C		381 (M <sup>+</sup> +1)
S-d-31	SG	S-c-101	nBuSO <sub>2</sub>	Et	2-Nap	C		425 (M <sup>+</sup> +1)
S-d-32	SA	S-d-31	nBuSO <sub>2</sub>	H	2-Nap	C		397 (M <sup>+</sup> +1)
S-d-33	SG	S-c-103	nBuSO <sub>2</sub>	Et	5-Ind	C		410 (M <sup>+</sup> +1)
S-d-34	SA	S-d-33	nBuSO <sub>2</sub>	H	5-Ind	C		385 (M <sup>+</sup> +1)
S-d-35	SG	S-c-105	nBuSO <sub>2</sub>	Et	1Me-5-1HIdz	C		425 (M <sup>+</sup> +1)
S-d-36	SA	S-d-35	nBuSO <sub>2</sub>	H	1Me-5-1HIdz	C		401 (M <sup>+</sup> +1)

[Reference Examples: Intermediates An-1 to An-5]

Synthesis of ethyl 3-[2-hydroxy-3-(naphthalen-2-yl)pyridin-5-yl]propionate

(Intermediate Ah-1)

A solution of the compound of Example P-42 (452 mg) in a mixture of ethyl acetate (5 ml) and methanol (2.5 ml) was added with 10% palladium/carbon (50 mg), and stirred at room temperature for 2 hours under hydrogen atmosphere. The

reaction mixture was filtered, and the solvent of the filtrate was evaporated under reduced pressure to obtain the title compound (Intermediate Ah-1, 321 mg). Mass (FAB): 322 (M<sup>+</sup>+1).

Synthesis of ethyl 3-[3-(naphthalen-2-yl)-2-(trifluoromethanesulfonyl)pyridin-5-yl]propionate (Intermediate An-1)

According to the procedure described in the synthesis method of Intermediate Aa-1, Intermediate Ah-1 (310 mg) and trifluoromethanesulfonic anhydride (170  $\mu$ l) were reacted and treated to obtain the title compound (Intermediate An-1, 355 mg). Mass (FAB): 454 (M<sup>+</sup>+1).

Typical examples of the reaction intermediates that can be obtained by reacting and treating corresponding starting compounds according to the method described above are shown below.

Intermediate An-2: ethyl 3-[3-(1H-indol-5-yl)-2-(trifluoromethanesulfonyl)pyridin-5-yl]propionate

Intermediate An-3: ethyl 3-[3-(1-methyl-1H-indol-5-yl)-2-(trifluoromethanesulfonyl)pyridin-5-yl]propionate

Intermediate An-4: ethyl 3-[3-(1H-indazol-5-yl)-2-(trifluoromethanesulfonyl)pyridin-5-yl]propionate

Intermediate An-5: ethyl 3-[3-(1-methyl-1H-indazol-5-yl)-2-(trifluoromethanesulfonyl)pyridin-5-yl]propionate

[Examples Cn-1 to Cn-45]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds according to the methods described in Examples Ca-1 and Ca-2 are shown in Table-Cn-1.

In the table, the substances mentioned in the column of "SM1" correspond to reaction intermediates, and the substances mentioned in the column of "SM2" correspond to the boronic acid reagent used in Example Ca-1. The boronic acid

reagents indicated with the symbols of “BRA (number)” in the columns of “SM2” are those mentioned in Table-Ba-1 and Table-Ba-2.

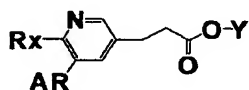


Table-Cn-1

Exp.	Rx	Y	AR	SM1	SM2	LCMS		
						method	RTime	Mass
Cn-1	Ph	Et	2-Nap	An-1	BRA14	D		382 (M <sup>+</sup> +1)
Cn-2	Ph	H	2-Nap	Cn-1	-	C		354 (M <sup>+</sup> +1)
Cn-3	Ph	Et	5-Ind	An-2	BRA14	C		371 (M <sup>+</sup> +1)
Cn-4	Ph	H	5-Ind	Cn-3	-	C		343 (M <sup>+</sup> +1)
Cn-5	Ph	Et	1Me-5-Ind	An-3	BRA14	C		385 (M <sup>+</sup> +1)
Cn-6	Ph	H	1Me-5-Ind	Cn-5	-	C		357 (M <sup>+</sup> +1)
Cn-7	Ph	Et	5-1HIdz	An-4	BRA14	C		372 (M <sup>+</sup> +1)
Cn-8	Ph	H	5-1HIdz	Cn-7	-	C		344 (M <sup>+</sup> +1)
Cn-9	Ph	Et	1Me-5-1HIdz	An-5	BRA14	C		386 (M <sup>+</sup> +1)
Cn-10	Ph	H	1Me-5-1HIdz	Cn-9	-	C		358 (M <sup>+</sup> +1)
Cn-11	4MeOPh	H	2-Nap	An-1	BRA19	C		384 (M <sup>+</sup> +1)
Cn-12	4MeOPh	H	5-Ind	An-2	BRA19	C		373 (M <sup>+</sup> +1)
Cn-13	4MeOPh	H	5-1HIdz	An-4	BRA19	C		374 (M <sup>+</sup> +1)
Cn-14	4MeOPh	H	1Me-5-1HIdz	An-5	BRA19	C		388 (M <sup>+</sup> +1)
Cn-15	3MeOPh	H	2-Nap	An-1	BRA37	C		384 (M <sup>+</sup> +1)
Cn-16	2MeOPh	H	2-Nap	An-1	BRA38	C		384 (M <sup>+</sup> +1)
Cn-17	2MeOPh	H	1Me-5-Ind	An-3	BRA38	C		387 (M <sup>+</sup> +1)
Cn-18	2MeOPh	H	1Me-5-1HIdz	An-5	BRA38	C		388 (M <sup>+</sup> +1)
Cn-19	2MePh	H	2-Nap	An-1	BRA59	C		368 (M <sup>+</sup> +1)
Cn-20	2MePh	H	1Me-5-Ind	An-3	BRA59	C		371 (M <sup>+</sup> +1)
Cn-21	2MePh	H	1Me-5-1HIdz	An-5	BRA59	C		372 (M <sup>+</sup> +1)
Cn-22	3MePh	H	2-Nap	An-1	BRA60	C		368 (M <sup>+</sup> +1)
Cn-23	3MePh	H	5-1HIdz	An-4	BRA60	C		358 (M <sup>+</sup> +1)
Cn-24	4MePh	H	2-Nap	An-1	BRA29	C		368 (M <sup>+</sup> +1)
Cn-25	4MePh	H	5-Ind	An-2	BRA29	C		357 (M <sup>+</sup> +1)
Cn-26	4MePh	H	1Me-5-Ind	An-3	BRA29	C		371 (M <sup>+</sup> +1)
Cn-27	4MePh	H	5-1HIdz	An-4	BRA29	C		358 (M <sup>+</sup> +1)
Cn-28	4MePh	H	1Me-5-1HIdz	An-5	BRA29	C		372 (M <sup>+</sup> +1)
Cn-29	4CF <sub>3</sub> Ph	H	5-Ind	An-2	BRA41	C		411 (M <sup>+</sup> +1)
Cn-30	4CF <sub>3</sub> Ph	H	5-1HIdz	An-4	BRA41	C		412 (M <sup>+</sup> +1)
Cn-31	4CF <sub>3</sub> Ph	H	1Me-5-1HIdz	An-5	BRA41	C		426 (M <sup>+</sup> +1)
Cn-32	4ClPh	H	5-Ind	An-2	BRA30	C		377 (M <sup>+</sup> +1)
Cn-33	4ClPh	H	1Me-5-Ind	An-3	BRA30	C		391 (M <sup>+</sup> +1)
Cn-34	4ClPh	H	1Me-5-1HIdz	An-5	BRA30	C		392 (M <sup>+</sup> +1)
Cn-35	2FPh	H	2-Nap	An-1	BRA32	C		372 (M <sup>+</sup> +1)
Cn-36	2FPh	H	1Me-5-Ind	An-3	BRA32	C		375 (M <sup>+</sup> +1)
Cn-37	2FPh	H	5-1HIdz	An-4	BRA32	C		362 (M <sup>+</sup> +1)
Cn-38	2FPh	H	1Me-5-1HIdz	An-5	BRA32	C		376 (M <sup>+</sup> +1)
Cn-39	3FPh	H	5-Ind	An-2	BRA33	C		361 (M <sup>+</sup> +1)
Cn-40	3FPh	H	5-1HIdz	An-4	BRA33	C		362 (M <sup>+</sup> +1)
Cn-41	3FPh	H	1Me-5-1HIdz	An-5	BRA33	C		376 (M <sup>+</sup> +1)
Cn-42	4FPh	H	2-Nap	An-1	BRA34	C		372 (M <sup>+</sup> +1)
Cn-43	4FPh	H	5-Ind	An-2	BRA34	C		361 (M <sup>+</sup> +1)
Cn-44	4FPh	H	5-1HIdz	An-4	BRA34	C		362 (M <sup>+</sup> +1)
Cn-45	4FPh	H	1Me-5-1HIdz	An-5	BRA34	C		376 (M <sup>+</sup> +1)

[Reference Examples: Intermediates Int. n-1 to Int. n-115]

Synthesis of methyl 3-(4-aminophenyl)propionate (Intermediate Int. n-1) (Synthesis

method NL)

A solution obtained beforehand by adding thionyl chloride (6.7 ml, WAKO) dropwise to methanol (50 ml) under ice cooling and mixing them was added dropwise with a solution of 4-aminohydrocinnamic acid (9.97 g, TCI) in methanol (50 ml) under ice cooling, stirred for 30 minutes, then warmed to room temperature, and further stirred for 16.5 hours. The reaction mixture was concentrated under reduced pressure, and then extracted with ethyl acetate (200 ml), and the organic layer was washed successively with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride and saturated brine. The organic layer was dried, and then the solvent was evaporated under reduced pressure to obtain the title compound (Intermediate Int. n-1, 13.13 g).

Synthesis of methyl 3-(4-amino-3-bromophenyl)propionate (Intermediate Int. n-2)  
(Synthesis method NK)

A solution of Intermediate Int. n-1 (9.93 g) in acetic acid (55 ml) was added with potassium bromide (6.60 g, WAKO) and sodium tungstenate(IV) dihydrate (18.23 g, WAKO), stirred for 5 minutes, then added dropwise with aqueous hydrogen peroxide (3.5 ml, WAKO) at 0°C over 5 minutes, warmed to room temperature, and then stirred for 1 hour. The reaction mixture was poured into 5% aqueous ammonia containing ice, thereby adjusted to pH of about 6, and then added with dichloromethane (200 ml) for extraction. The organic layer was washed successively with saturated aqueous ammonium chloride, saturated aqueous sodium hydrogencarbonate and saturated brine, and then dried, and the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 3:1) to obtain the title compound (Intermediate Int. n-2, 3.07 g).

Synthesis of methyl 3-(4-benzylamino-3-bromophenyl)propionate (Intermediate Int. n-3) (Synthesis method NC1)



A solution of Intermediate Int. n-2 (10.97 g) in methanol (30 ml) was added with benzaldehyde (5.25 ml, TCI) and anhydrous sodium sulfate (6.49 g, WAKO), and stirred at 60°C for 13 hours. The reaction mixture was added with sodium cyanotrihydridoborate (2.73 g, WAKO), and further stirred for 5 hours. The reaction mixture was concentrated under reduced pressure, and then extracted with dichloromethane (150 ml), and the organic layer was washed with saturated brine, and dried. Then, the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 4:1) to obtain the title compound (Intermediate Int. n-3, 13.45 g).

Synthesis of methyl 3-[3-bromo-(4-fluorobenzylamino)phenyl]propionate (Intermediate Int. n-4) (Synthesis method NC2)

A solution of Intermediate Int. n-2 (5.80 g) in dichloromethane (100 ml) was added with p-fluorobenzaldehyde (2.83 ml, TCI), sodium triacetoxyborohydride (7.14 g, Ald) and acetic acid (1.4 ml), and stirred at room temperature for 19 hours. The reaction mixture was extracted with dichloromethane (300 ml), and the organic layer was washed with saturated brine, and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 4:1) to obtain the title compound (Intermediate Int. n-4, 7.51 g).

Synthesis of methyl 3-[4-amino-3-(naphthalen-2-yl)phenyl]propionate (Intermediate Int. n-7) (Synthesis method ND1)

A solution of the compound of Example N-a-1 (3.01 g) in a mixture of methanol (40 ml) and THF (20 ml) was added with 10% palladium/carbon (410.3 mg, Merck) and one drop of concentrated hydrochloric acid, and stirred at room temperature for 5 hours under hydrogen atmosphere. The reaction mixture was filtered, and the solvent of the filtrate was evaporated under reduced pressure. The residue was added with ethyl acetate (200 ml), and washed successively with

saturated aqueous sodium hydrogencarbonate and saturated brine, and then dried, and the solvent was evaporated under reduced pressure to obtain the title compound (Intermediate Int. n-7, 2.58 g).

Synthesis of methyl 3-[3-nitro-4-(piperazin-1-yl)phenyl]acrylate (Intermediate Int. n-19) (Synthesis method NJ)

A solution of methoxycarbonylmethyl(triphenyl)phosphonium bromide (1.1 g, TCI) in THF (12.5 ml) was added with sodium hydride (115 mg, WAKO) under ice cooling, warmed to room temperature, then added dropwise with a solution of 3-nitro-4-(piperazin-1-yl)benzaldehyde (550.6 mg, MAYB) in THF (12.5 ml), and stirred at the same temperature for 16.5 hours. The reaction mixture was poured into brine (40 ml), and extracted with ethyl acetate (100 ml). The organic layer was washed successively with saturated aqueous sodium hydrogencarbonate and saturated brine, and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography (hexane:ethyl acetate = 5:1) to obtain the title compound (Intermediate Int. n-19, 511 mg).

Synthesis of methyl 3-[3-amino-4-(piperazin-1-yl)phenyl]propionate (Intermediate Int. n-20) (Synthesis method ND1)

According to the procedure described in the synthesis method of Intermediate Int. n-7 (Synthesis method ND1) provided that the reaction was carried out in ethyl acetate for 13 hours, Intermediate Int. n-19 (505 mg) and 10% palladium/carbon (50 mg) were reacted and treated to obtain the title compound (Intermediate Int. n-20, 658.9 mg).

Synthesis of methyl 3-[3-bromo-4-(piperazin-1-yl)phenyl]propionate (Intermediate Int. n-21) (Synthesis method NI)

A solution of hydrobromic acid (570  $\mu$ l) in methanol (2.3 ml) was added dropwise with a solution of Intermediate Int. n-20 (235 mg) in methanol (2.3 ml) over 10 minutes under ice cooling. This reaction mixture was added with an

aqueous solution (250  $\mu$ l) of sodium nitrite (69 mg, WAKO). The reaction mixture was added dropwise with an aqueous solution (2.3 ml) of copper(II) bromide (222 mg, WAKO) heated to 50°C over 15 minutes, stirred for 4 hours at the same temperature, and then further stirred at room temperature for 12.5 hours. The reaction mixture was poured into aqueous sodium hydrogencarbonate (20 ml), and extracted with ethyl acetate (40 ml). The organic layer was washed with saturated brine, and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography (hexane:ethyl acetate = 4:1) to obtain the title compound (Intermediate Int. n-21, 89 mg).

Synthesis of methyl 4-fluoro-3-bromocinnamate (Intermediate Int. n-25) (Synthesis method NL)

According to the procedure described in the synthesis method of Intermediate Int. n-1 (Synthesis method NL) provided that the reaction was carried out for 1 hour, 3-bromo-4-fluorocinnamic acid (3.30 g, LANC) and thionyl chloride (1.5 ml) were reacted and treated to obtain the title compound (Intermediate Int. n-25, 3.47 g)

Synthesis of methyl 3-[3-bromo-4-(piperidin-1-yl)phenyl]cinnamate (Intermediate Int. n-26) (Synthesis method NG)

A solution of Intermediate Int. n-25 (136.4 mg) in DMSO (5 ml) was added with potassium carbonate (109.8 mg) and piperidine (84.8  $\mu$ l, TCI), and stirred at 90°C for 15 hours. The reaction mixture was extracted with ethyl acetate (50 ml), and then the organic layer was washed successively with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride and saturated brine, and dried. Then, the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:isopropyl ether = 6:1) to obtain the title compound (Intermediate Int. n-26, 120.4 mg).

Synthesis of methyl 3-[3-bromo-4-(piperidin-1-yl)phenyl]propionate (Intermediate

Int. n-27) (Synthesis method ND2)

According to a procedure described in literature [D.J. Hart et al., *Journal of Organic Chemistry* (J. Org. Chem.), 1987, vol. 52, p.4665], a solution of Intermediate Int. n-26 (690.6 mg) in dimethoxyethane (100 ml) was added with p-toluenesulfonhydrazide (2.97g, TCI), and refluxed by heating at 110°C. Then, the reaction mixture was added dropwise with an aqueous solution (100 ml) of sodium acetate (2.85 g, WAKO) over 2 hours, and further stirred for 1 hour. The reaction mixture was extracted with dichloromethane (450 ml), and the organic layer was washed with water, and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 6:1) to obtain the title compound (Intermediate Int. n-27, 648.2 mg).

Synthesis of 3-bromo-(4-imidazol-1-yl)benzaldehyde (Intermediate Int. n-32)  
(Synthesis method NG)

According to the procedure described in the synthesis method of Intermediate Int. n-26 (Synthesis method NG) provided that the reaction was performed for 20 hours, and the column chromatography was performed with chloroform:methanol = 100:1, 3-bromo-4-fluorobenzaldehyde (1.246 g, TCI), potassium carbonate (825.1 mg) and imidazole (444 mg, TCI) were reacted and treated to obtain the title compound (Intermediate Int. n-32, 986.1 mg).

Synthesis of ethyl 3-[3-bromo-(4-imidazol-1-yl)phenyl]acrylate (Intermediate Int. n-33) (Synthesis method NJ)

A solution of Intermediate Int. n-32 (986.1 mg) and ethyl diethylphosphonoacetate (705  $\mu$ l) in 1,2-dimethoxyethane (8 ml) was added with 60% sodium hydride (180.2 mg) under ice cooling, stirred for 10 minutes, then warmed to room temperature, and stirred for 1 hour. The reaction mixture was added with dichloromethane (60 ml) for extraction, and the organic layer was

washed with saturated brine, and dried. Then, the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, dichloromethane:methanol = 100:1) to obtain the title compound (Intermediate Int. n-33, 1.00 g).

Synthesis of methyl 3-(4-cyclopentylaminophenyl)propionate (Intermediate Int. n-38) (Synthesis method NC1)

According to the procedure described in the synthesis method of Intermediate Int. n-3 provided that the reaction was carried out for 6 hours, Intermediate Int. n-1 (1.03 g), cyclopentanone (450  $\mu$ l, TCI), sodium triacetoxyborohydride (1.56 g) and acetic acid (350  $\mu$ l) were reacted and treated to obtain the title compound (Intermediate Int. n-37, 1.21 g).

Synthesis of methyl 3-(4-cyclopentylamino-3,5-dibromophenyl)propionate (Intermediate Int. n-39) (Synthesis method NK)

A solution of Intermediate Int. n-37 (1.21 g) in acetonitrile was warmed to 35°C, then added with N-bromosuccinimide (2.44 g, TCI), and stirred for 1 hour. The reaction mixture was concentrated under reduced pressure, then added with ethyl acetate (150 ml), washed successively with aqueous sodium thiosulfate, saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride and saturated brine, and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 6:1) to obtain the title compound (Intermediate Int. n-38, 1.41 g).

Synthesis of 2-bromopyridine-5-carbaldehyde (Intermediate Int. n-44) (Synthesis method NM)

According to a procedure described in literature (Xin Wang et al., Tetrahedron. Lett., 2000, vol. 41, p.4335], a solution of 2,5-dibromopyridine (3.17 g) in anhydrous diethyl ether (140 ml) was added dropwise with a 1.6 M solution of n-

butyl lithium in hexane (11 ml) with cooling at -78°C under argon gas atmosphere over 5 minutes, and stirred for 20 minutes. This reaction mixture was added dropwise with dehydrated DMF (1 ml) over 3 minutes, stirred for 30 minutes, then warmed to room temperature, and further stirred for 1 hour. The reaction mixture was added with water (20 ml), and extracted with ethyl acetate (30 ml x 2). The organic layer was washed with saturated brine, and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 6:1) to obtain the title compound (Intermediate Int. n-44, 1.46 g).

Synthesis of ethyl 3-(2-bromopyridin-5-yl)acrylate (Intermediate Int. n-45)  
(Synthesis method NJ)

According to the procedure described in the synthesis method of Intermediate n-7 provided that the reaction was carried out for 15 minutes, Intermediate Int. n-44 (1.45 g), ethyl diethylphosphonoacetate (2.1 ml) and 60% sodium hydride (355 mg) were reacted and treated to obtain the title compound (Intermediate Int. n-45, 1.87 g).

Synthesis of ethyl 3-[2-(piperidin-1-yl)pyridin-5-yl]acrylate (Intermediate Int. n-46)  
(Synthesis method NG)

Intermediate Int. n-45 (565.7 mg) was added with potassium carbonate (286.4 mg) and piperidine (3 ml), and stirred at 90°C for 21 hours. The reaction mixture was added with ethyl acetate (50 ml), washed successively with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride and saturated brine, and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 4:1) to obtain the title compound (Intermediate Int. n-46, 165.9 mg).

Synthesis of ethyl 3-[2-(piperidin-1-yl)pyridin-5-yl]propionate (Intermediate Int. n-

47) (Synthesis method ND1)

According to the procedure described in the synthesis method of Intermediate Int. n-7 with the modifications that the reaction was carried out for 1 hour, and the purification was performed by column chromatography (Quad, hexane:ethyl acetate = 6:1), Intermediate Int. n-46 (392 mg) and 10% palladium/carbon (30 mg) were reacted and treated to obtain the title compound (Intermediate Int. n-47; 246 mg).

Synthesis of ethyl 3-[3-bromo-2-(piperidin-1-yl)pyridin-3-yl]propionate (Intermediate Int. n-48) (Synthesis method NK2)

A solution of Intermediate Int. n-47 (242 mg) in acetonitrile was added with bromine (84  $\mu$ l), and stirred at 40°C for 1 hour. The reaction mixture was concentrated under reduced pressure, then added with ethyl acetate (50 ml), washed successively with aqueous sodium thiosulfate, saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride and saturated brine, and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 6:1) to obtain the title compound (Intermediate Int. n-48, 224 mg).

Synthesis of 2-benzylaminopyridine-5-carbaldehyde (Intermediate Int. n-59) (Synthesis method NG)

Intermediate Int. n-44 (102.0 mg) was added with benzylamine (1 ml, TCI), and stirred at 120°C for 39 hours. The reaction mixture was added with ethyl acetate (50 ml), washed successively with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride and saturated brine, and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 4:1) to obtain the title compound (Intermediate Int. n-59, 58.3 mg).

Synthesis of 2-benzylamino-3-bromopyridine-5-carbaldehyde (Intermediate Int. n-

60) (Synthesis method NK)

A solution of Intermediate Int. n-59 (56.8 mg) in acetonitrile was added with N-bromosuccinimide (134 mg), and stirred at room temperature for 14 hours. The reaction mixture was concentrated under reduced pressure, then added with ethyl acetate (50 ml), washed successively with aqueous sodium thiosulfate, saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride and saturated brine, and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 4:1) to obtain the title compound (Intermediate Int. n-60, 50 mg).

Synthesis of ethyl 3-(2-benzylamino-3-bromopyridin-5-yl)acrylate (Intermediate Int. n-61) (Synthesis method NJ)

According to the procedure described in the synthesis method of Intermediate Int. n-7 provided that the reaction was carried out for 30 minutes, Intermediate Int. n-60 (49.1 g), ethyl diethylphosphonoacetate (92  $\mu$ l) and 60% sodium hydride (30 mg) were reacted and treated to obtain the title compound (Intermediate Int. n-61, 28 mg).

Synthesis of ethyl 3-(2-benzylamino-3-bromopyridin-5-yl)propionate (Intermediate Int. n-62) (Synthesis method ND2)

According to the procedure described in the synthesis method of Intermediate Int. n-27 provided that the reaction was carried out for 4 hours, Intermediate Int. n-60 (49.1 mg), p-toluenesulfonhydrazide (320.6 mg) and sodium acetate (412.4 mg) were reacted and treated to obtain the title compound (Intermediate Int. n-62, 38.9 mg).

Synthesis of methyl 3-(4-amino-3-bromo-5-nitrophenyl)propionate (Intermediate Int. n-76) (Synthesis method NM)

A solution obtained by adding potassium nitrate (1.10 g) to a solution of










Intermediate Int. n-2 (2.57 g) in acetic anhydride (20 ml) under ice cooling and stirring them for 10 minutes was added dropwise with concentrated sulfuric acid (700  $\mu$ l) over 10 minutes. The reaction mixture was stirred for 10 minutes at the same temperature, then warmed to room temperature, and further stirred for 30 minutes. The reaction mixture was poured into 1 N aqueous sodium hydroxide (250 ml) containing ice, and extracted with isopropyl ether (200 ml x 2). The organic layer was washed successively with saturated aqueous sodium hydrogencarbonate and saturated brine, and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 3:1) to obtain the title compound (Intermediate Int. n-76, 0.72 g).

Typical examples of the intermediates for synthesizing the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification are shown in Table-Int. N-1 to Table-Int. N-8. In the tables, the intermediate numbers "Int. n-(number)" are mentioned in the columns indicated as "Exp.". In the tables, used methods among the aforementioned synthesis methods are shown in the columns of "Syn" with symbols, the starting compounds 1 are mentioned in the columns of "SM1", and the starting compounds 2 are mentioned in the columns of "SM2". Further, the compounds indicated as "Single" in the columns of "Single or Double" in the tables are compound in which two of the carbon atoms binding the benzene ring and carbonyl group in the compounds are bound with a single bond, and those indicated as "Double" in the same are compounds in which two of the carbon atoms binding the benzene ring and carbonyl group in the compounds are bound with a double bond. The aldehydes and ketones used for the synthesis of the compounds are mentioned in Table-Carb, and amines used for the same are mentioned in Table-AMN.

Table-Carb

Reagent	Aldehyde or Ketone	Manufacture
CHO1	HCHO	WAKO
CHO2	CH <sub>3</sub> CHO	Aldrich
CHO3	CH <sub>3</sub> CH <sub>2</sub> CHO	TCI
CHO4	nPrCHO	TCI
CHO5	Acetone	WAKO
CHO6	nBuCHO	TCI
CHO7	iPrCHO	TCI
CHO8	BnCHO	TCI
CHO9	4FBnCHO	TCI
CHO10	2FBnCHO	TCI
CHO11	3FBnCHO	TCI
CHO12	2ClBnCHO	TCI
CHO13	2BrBnCHO	TCI
CHO14	2,3DFBnCHO	TCI
CHO15	3,4DFBnCHO	TCI
CHO16	4PhBnCHO	TCI
CHO17	2CF <sub>3</sub> BnCHO	TCI
CHO18	2,3DCIBnCHO	TCI
CHO19	2-ThiopheneCHO(2-TFCHO)	TCI
CHO20	3-ThiopheneCHO(3-TFCHO)	TCI
CHO21	2-FuranCHO(2-FRCHO)	TCI
CHO22	Cyclopentanone	TCI
CHO23	Cyclohexanone	TCI
CHO24	2(Me)cHexanone	TCI
CHO25	2-Indanone	Aldrich

Table-AMN

Reagent	Amine	Manufacture
AMN1		TCI
AMN2		TCI
AMN3		TCI
AMN4		TCI
AMN5		TCI
AMN6		TCI
AMN7		TCI
AMN8	EtMeNH	Aldrich
AMN9	Et <sub>2</sub> NH	Aldrich
AMN10	nPrMeNH	Aldrich
AMN11	iPrMeNH	Aldrich
AMN12	nBuMeNH	Aldrich
AMN13	nBuEtNH	Aldrich
AMN14	iBuMeNH	Aldrich
AMN15	4MeBnNH <sub>2</sub>	Aldrich
AMN16	3MeBnNH <sub>2</sub>	Aldrich
AMN17	2MeBnNH <sub>2</sub>	Aldrich
AMN18	4FBnNH <sub>2</sub>	Aldrich
AMN19	3FBnNH <sub>2</sub>	Aldrich
AMN20	2FBnNH <sub>2</sub>	Aldrich
AMN21	3MeOBnNH <sub>2</sub>	Aldrich
AMN22	4MeOBnNH <sub>2</sub>	Aldrich
AMN23	2MeOBnNH <sub>2</sub>	Aldrich
AMN24	4CF <sub>3</sub> BnNH <sub>2</sub>	Aldrich
AMN25	2EtOBnNH <sub>2</sub>	Aldrich
AMN26	3iPrOBnNH <sub>2</sub>	Sigma-Aldrich
AMN27	3,5DFBnNH <sub>2</sub>	Aldrich

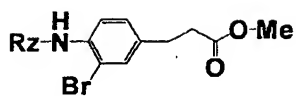


Table-Int.N-1

Exp.	Syn	SM1	SM2	Rz	LCMS		
					method	RTime	Mass
Int.n-5	NC2	Int.n-2	CHO10	2FBn	C		366 (M <sup>+</sup> )
Int.n-6	NC1	Int.n-2	CHO11	3FBn	C		366 (M <sup>+</sup> )
Int.n-12	NC2	Int.n-2	CHO2	Et	C		386 (M <sup>+</sup> )
Int.n-13	NC2	Int.n-2	CHO3	nPr	D	5.02	300 (M <sup>+</sup> )
Int.n-14	NC2	Int.n-2	CHO5	iPr	D	5.38	341(M <sup>+</sup> )
Int.n-15	NC2	Int.n-2	CHO7	iBu	D	5.50	400 (M <sup>+</sup> )
Int.n-16	NC2	Int.n-2	CHO22	cPen	C		326 (M <sup>+</sup> )
Int.n-17	NC2	Int.n-2	CHO23	cHex	C		340(M <sup>+</sup> )
Int.n-18	NC2	Int.n-2	CHO24	2(Me)cHex	C		354 (M <sup>+</sup> )

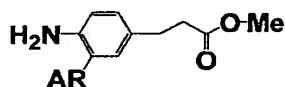


Table-Int.N-2

Exp.	Syn	SM1	AR	LCMS		
				method	RTime	Mass
Int.n-8	ND1	N-a-3	5-Ind	C		295 (M <sup>+</sup> +1)
Int.n-9	ND1	N-a-5	1Me-5-Ind	C		309 (M <sup>+</sup> +1)
Int.n-10	ND1	N-a-7	5-1Idz	C		296 (M <sup>+</sup> +1)
Int.n-11	ND1	N-a-9	1Me-5-1HIdz	C		310(M <sup>+</sup> +1)

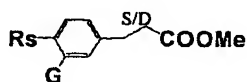


Table-Int.N-3

Exp.	Syn.	Rs	G	Single or Double	LCMS		
					method	RTime	Mass
Int.n-22	NJ		NO2	Double	A	3.91	293 (M <sup>+</sup> +1)
Int.n-23	ND1		NH2	Single	A	2.97	265(M <sup>+</sup> +1)
Int.n-24	NI		Br	Single	A	4.31	328 (M <sup>+</sup> )

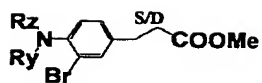


Table-Int.N-4

Exp.	Syn	SM1	RzRyN	Single or Double	LCMS		
					method	RTime	Mass
Int.n-28	NC2	Int.n-25		Double	A	6.54	338(M <sup>+</sup> )
Int.n-29	NC1	Int.n-28		Single	A	6.01	342 (M <sup>+</sup> +1)
Int.n-30	NC2	Int.n-25		Double	A	6.29	340(M <sup>+</sup> +1)
Int.n-31	NC1	Int.n-30		Single	A	6.12	342 (M <sup>+</sup> +1)
Int.n-33	NC2	Int.n-32		Double	C		307 (M <sup>+</sup> )
Int.n-35	NC2	Int.n-32		Double	C		306 (M <sup>+</sup> )
Int.n-36	NC2	Int.n-25		Double	A	5.60	310 (M <sup>+</sup> )
Int.n-37	NC2	Int.n-32		Double	C		326 (M <sup>+</sup> )

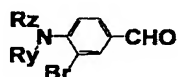


Table-Int.N-5

Exp.	Syn	SM1	RzRyN	LCMS		
				method	RTime	Mass
Int.n-32	NC2	Int.n-25		C		351 (M <sup>+</sup> )
Int.n-34	NC2	Int.n-25		C		250 (M <sup>+</sup> )

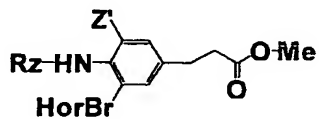


Table-Int.N-6

Exp.	Syn	SM1	SM2	Rz	Z'	H or Br	LCMS		
							method	RTime	Mass
Int.n-40	NC1	Int.n-1	CHO3	nPr	H	H	C		222 (M <sup>+</sup> +1)
Int.n-41	NK	Int.n-40		nPr	Br	Br	C		380 (M <sup>+</sup> +1)
Int.n-42	NC1	Int.n-1	CHO5	iPr	H	H	C		222 (M <sup>+</sup> +1)
Int.n-43	NK	Int.n-42		iPr	Br	Br	C		380 (M <sup>+</sup> +1)

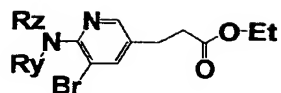



Table-Int.N-7

Exp.	Syn	RzRyN		LCMS		
				method	RTime	Mass
Int.n-48	NG	cHex		C		327 (M <sup>+</sup> )
Int.n-49	NG	cPen		C		313(M <sup>+</sup> )
Int.n-50	NG	4(Me)cHex		C		341 (M <sup>+</sup> )
Int.n-51	NG			C		343 (M <sup>+</sup> )
Int.n-52	NG	cHep		C		355 (M <sup>+</sup> )
Exp.	Syn	Rz	Ry	LCMS		
				method	RTime	Mass
Int.n-53	NG	Et	Me	C		301 (M <sup>+</sup> )
Int.n-54	NG	Et	Et	C		315 (M <sup>+</sup> )
Int.n-55	NG	nPr	Me	C		315 (M <sup>+</sup> )
Int.n-56	NG	iPr	Me	C		315 (M <sup>+</sup> )
Int.n-57	NG	nBu	Me	C		329 (M <sup>+</sup> )
Int.n-58	NG	iBu	Me	C		329 (M <sup>+</sup> )
Int.n-63	NG	4MeBn	H	C		363 (M <sup>+</sup> )
Int.n-64	NG	3MeBn	H	C		363 (M <sup>+</sup> )
Int.n-65	NG	2MeBn	H	C		363 (M <sup>+</sup> )
Int.n-66	NG	4FBn	H	C		368 (M <sup>+</sup> +1)
Int.n-67	NG	3FBn	H	C		368 (M <sup>+</sup> +1)
Int.n-68	NG	2FBn	H	C		368 (M <sup>+</sup> +1)
Int.n-69	NG	4MeOPh	H	C		365 (M <sup>+</sup> )
Int.n-70	NG	3MeOPh	H	C		365 (M <sup>+</sup> )
Int.n-71	NG	2MeOPh	H	C		365 (M <sup>+</sup> )
Int.n-72	NG	4CF3Ph	H	C		403 (M <sup>+</sup> )
Int.n-73	NG	2EtOPh	H	C		380 (M <sup>+</sup> +1)
Int.n-74	NG	3iPrOPh	H	C		393 (M <sup>+</sup> )
Int.n-75	NG	3,5DFPh	H	C		372 (M <sup>+</sup> +1)

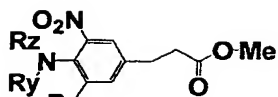



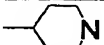



Table-Int.N-8

Exp.	Syn	SM1	SM2	Rz	Ry	LCMS		
						method	RTime	Mass
Intn-77	NC2	Intn-76	CHO22	cPen	H	C		371 (M <sup>+</sup> )
Intn-78	NC2	Intn-76	CHO3	nPr	H	C		345 (M <sup>+</sup> )
Intn-79	NC2	Intn-76	CHO5	iPr	H	C		345 (M <sup>+</sup> )
Intn-80	NC2	Intn-76	CHO25	2-Indane	H	C		419 (M <sup>+</sup> )
Intn-81	NC2	Intn-76	CHO23	cHex	H	C		385 (M <sup>+</sup> )
Intn-82	NC2	Intn-76	CHO24	2(Me)cHex	H	C		399 (M <sup>+</sup> )
Intn-83	NC1	Intn-77	CHO1	cPen	Me	C		385 (M <sup>+</sup> )
Intn-84	NC1	Intn-78	CHO1	nPr	Me	C		359(M <sup>+</sup> )
Intn-85	NC1	Intn-79	CHO1	iPr	Me	C		359(M <sup>+</sup> )
Intn-86	NC1	Intn-80	CHO1	2-Indane	Me	C		433 (M <sup>+</sup> )
Intn-87	NC1	Intn-81	CHO1	cHex	Me	C		399 (M <sup>+</sup> )
Intn-88	NC1	Intn-82	CHO1	2(Me)cHex	Me	C		413 (M <sup>+</sup> )
Intn-89	NC1	Intn-76	CHO8	Bn	H	C		393 (M <sup>+</sup> )
Intn-90	NC1	Intn-76	CHO9	4FBn	H	C		411 (M <sup>+</sup> )
Intn-91	NC2	Intn-76	CHO10	2FBn	H	C		411 (M <sup>+</sup> )
Intn-92	NC2	Intn-76	CHO11	3FBn	H	C		411 (M <sup>+</sup> )
Intn-93	NC2	Intn-76	CHO14	2,3DFBn	H	C		429 (M <sup>+</sup> )
Intn-94	NC2	Intn-76	CHO15	3,4DFBn	H	C		429 (M <sup>+</sup> )
Intn-95	NC2	Intn-76	CHO16	4PhBn	H	C		469 (M <sup>+</sup> )
Intn-96	NC2	Intn-76	CHO17	2CF3Bn	H	C		461 (M <sup>+</sup> )
Intn-97	NC2	Intn-76	CHO19	2-TF	H	C		399 (M <sup>+</sup> )
Intn-98	NC2	Intn-76	CHO20	3-TF	H	C		399 (M <sup>+</sup> )
Intn-99	NC2	Intn-76	CHO21	2-FR	H	C		383 (M <sup>+</sup> )
Intn-100	NC1	Intn-89	CHO1	Bn	Me	C		407 (M <sup>+</sup> )
Intn-101	NC1	Intn-90	CHO1	4FBn	Me	C		428 (M <sup>+</sup> )
Intn-102	NC1	Intn-91	CHO1	2FBn	Me	C		425 (M <sup>+</sup> )
Intn-103	NC1	Intn-92	CHO1	3FBn	Me	C		425 (M <sup>+</sup> )
Intn-104	NC1	Intn-93	CHO1	2,3DFBn	Me	C		443 (M <sup>+</sup> )
Intn-105	NC1	Intn-94	CHO1	3,4DFBn	Me	C		443 (M <sup>+</sup> )
Intn-106	NC1	Intn-95	CHO1	4PhBn	Me	C		483 (M <sup>+</sup> )
Intn-107	NC1	Intn-96	CHO1	2CF3Bn	Me	C		475 (M <sup>+</sup> )
Intn-108	NC1	Intn-97	CHO1	2-TF	Me	C		413 (M <sup>+</sup> )
Intn-109	NC1	Intn-98	CHO1	3-TF	Me	C		413 (M <sup>+</sup> )
Intn-110	NC1	Intn-99	CHO1	2-FR	Me	C		397 (M <sup>+</sup> )
Exp.	Syn	SM1	SM2	RzRyN	LCMS			
					method	RTime	Mass	
Intn-111	NM	Intn-21			C		357 (M <sup>+</sup> )	
Intn-112	NM	Intn-24			C		373 (M <sup>+</sup> )	
Intn-113	NM	Intn-27			C		371 (M <sup>+</sup> )	
Intn-114	NM	Intn-29			C		385 (M <sup>+</sup> )	
Intn-115	NM	Intn-31			C		385 (M <sup>+</sup> )	

[Example N-a-1]

Synthesis of methyl 3-[4-benzylamino-3-(naphthalen-2-yl)phenyl]propionate  
(Compound No. N-a-1) (Synthesis method NB1)

A solution of Intermediate n-3 (8.18 g) in toluene (60 ml) was added with 2-naphthaleneboronic acid (5.04 g, TCI), 2 M aqueous sodium carbonate (21.6 ml), methanol (24 ml) and tetrakis(triphenylphosphine) palladium(0) (henceforth abbreviated as "(Ph<sub>3</sub>P)<sub>4</sub>Pd", 1.94 g, Nacalai Tesque), and stirred at 90°C for 15 hours. The reaction mixture was added with ethyl acetate (300 ml), and washed successively with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride, and saturated brine. The organic layer was dried, and then the solvent was evaporated under reduced pressure. The residue was purified by flash column chromatography (hexane:ethyl acetate = 3:1) to obtain the title compound (Compound No. N-a-1, 5.70 g).

[Example N-a-2]

Synthesis of 3-[4-benzylamino-3-(naphthalen-2-yl)phenyl]propionic acid (Compound No. N-a-2) (Synthesis method NA)

A solution of the compound of Example N-a-1 (51 mg) in methanol (5.0 ml) was added with 2 N aqueous sodium hydroxide (130  $\mu$ l), and stirred at 60°C for 2 hours. The reaction mixture was concentrated under reduced pressure, then neutralized with 5% aqueous hydrochloric acid under ice cooling, and then extracted with ethyl acetate (30 ml). The organic layer was washed with saturated brine, and dried, and then the solvent was evaporated under reduced pressure to obtain the title compound (Compound No. N-a-2, 47 mg).

[Example N-a-25]

Synthesis of methyl 3-[4-(N-benzyl-N-methylamino)-3-(naphthalen-2-yl)phenyl]propionate (Compound No. N-a-25) (Synthesis method NC1)

According to the procedure described in the synthesis method of



Intermediate n-3 provided that the reaction was carried out for 5 hours, the compound of Example N-a-1 (234.2 mg), 30% aqueous solution of formaldehyde (208.8  $\mu$ l, WAKO) and sodium cyanotrihydridoborate (140.9 mg) were reacted and treated to obtain the title compound (Compound No. N-a-25, 176.3 mg).

[Example N-A-137]

Synthesis of methyl 3-{3-(1-methyl-1H-indol-5-yl)-4-[N-(1-phenylethyl)amino]phenyl}propionate (Compound No. N-a-137) (Synthesis method NE1)

According to a procedure described in literature [Shin-Shyong Tseng et al., Journal of Organic Chemistry (J. Org. Chem.), 1979, vol. 44, p.4113], a solution of Intermediate n-9 (630.7 mg) in methylene chloride (10 ml) was added with triethylamine (405  $\mu$ l, Kokusan Chemical), cooled to -78°C, then added dropwise with trifluoromethanesulfonyl chloride (426  $\mu$ l, TCI), and stirred for 1.5 hours. The reaction mixture was poured into ice water (10 ml), and added with dichloromethane (30 ml) for extraction. The organic layer was washed with saturated brine, and dried, and then the solvent was evaporated under reduced pressure to obtain a crude product. A solution of the obtained crude product in DMF (15 ml) was added with potassium carbonate (394.2 mg) and (1-bromoethyl)benzene (386.4  $\mu$ l, TCI), and stirred at room temperature for 13 hours. The reaction mixture was extracted with ethyl acetate (100 ml), and the organic layer was washed successively with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride and saturated brine, and dried. Then, the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 4:1) to obtain the title compound (Compound No. N-a-137, 310.3 mg).

[Example N-a-141]

Synthesis of methyl 3-[3-(1-methyl-1H-indol-5-yl)-4-{N-[2-(4-

fluorophenyl)ethyl]amino}phenyl]propionate (Compound No. N-a-141) (Synthesis method NE2)

A solution of Intermediate n-9 (210.1 mg) in methylene chloride (10 ml) was added with triethylamine (135  $\mu$ l, Kokusan Chemical), cooled to  $-78^{\circ}\text{C}$ , then added dropwise with trifluoromethanesulfonyl chloride (143  $\mu$ l, TCI), and stirred for 1.5 hours. The reaction mixture was poured into ice water (10 ml), and added with dichloromethane (15 ml) for extraction. The organic layer was washed with saturated brine, and dried, and then the solvent was evaporated under reduced pressure to obtain a crude product. A solution of the obtained crude product in anhydrous DMF (15 ml) was added with triphenylphosphine (485.9 g, WAKO), di-*t*-butyl azodicarboxylate (299.8 mg, Ald) and 4-fluorophenylethyl alcohol (357  $\mu$ l, TCI), and stirred at room temperature for 12 hours. The reaction mixture was added with water (10 ml) and ethyl acetate (10 ml) for extraction, and the organic layer was washed successively with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride, and saturated brine, and dried. Then, the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 4:1) to obtain the title compound (Compound No. N-a-141, 63.5 mg).

[Example N-a-143]

Synthesis of methyl 3-[4-(N-acetyl-N-benzylamino)-3-(1-methyl-1H-indol-5-yl)phenyl]propionate (Compound No. N-a-143) (Synthesis method NF)

A solution of Compound No. N-a-5 (32 mg) in methylene chloride (3 ml) was added with pyridine (49.6  $\mu$ l, TCI) and acetyl chloride (50  $\mu$ l, TCI), and stirred for 13 hours. The reaction mixture was added with water (1 ml), and the solvent was evaporated. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 4:1) to obtain the title compound (Compound No. N-a-143, 20.3 mg).

[Example N-a-153]

Synthesis of methyl 3-[4-benzoylamino-3-(1-methyl-1H-indol-5-yl)phenyl]propionate  
(Compound No. N-a-153) (Synthesis method NF)

According to the procedure described in the synthesis method of the compound of Example N-a-143 provided that the reaction was carried out for 16 hours, Intermediate Int. n-9 (26.5 mg), pyridine (23.8  $\mu$  l) and benzoyl chloride (30  $\mu$  l, WAKO) were reacted and treated to obtain the title compound (Compound No. N-a-153, 18.4 mg).

[Examples N-a-1 to N-a-166]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification including the examples described above are shown in Table-N-A-1 to Table-N-A-4. In the tables, the compound numbers are mentioned in the columns indicated as "Exp.". In the tables, used methods among the aforementioned synthesis methods are shown in the columns of "Syn" with symbols, the starting compounds 1 are mentioned in the columns of "SM1", and the starting compounds 2 are mentioned in the columns of "SM2".

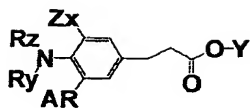


Table-N-A-1

Exp.	Syn	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									method	RTime	Mass
N-a-1	NB1	Int.n-3	BRN1	Bn	H	Me	H	2-Nap	C		396 (M <sup>+</sup> +1)
N-a-2	NA	N-a-1		Bn	H	H	H	2-Nap	C		382 (M <sup>+</sup> +1)
N-a-3	NB1	Int.n-3	BRN2	Bn	H	Me	H	5-1Ind	C		385 (M <sup>+</sup> +1)
N-a-4	NA	N-a-3		Bn	H	H	H	5-1Ind	C		371 (M <sup>+</sup> +1)
N-a-5	NB1	Int.n-3	BRN3	Bn	H	Me	H	1Me-5-Ind	C		399 (M <sup>+</sup> +1)
N-a-6	NA	N-a-5		Bn	H	H	H	1Me-5-Ind	C		385 (M <sup>+</sup> +1)
N-a-7	NB1	Int.n-3	BRN4	Bn	H	Me	H	1Et-5-Ind	C		413 (M <sup>+</sup> +1)
N-a-8	NA	N-a-7		Bn	H	H	H	1Et-5-Ind	C		399 (M <sup>+</sup> +1)
N-a-9	NB1	Int.n-3	BRN5	Bn	H	Me	H	5-1HIdz	C		386 (M <sup>+</sup> +1)
N-a-10	NA	N-a-9		Bn	H	H	H	5-1HIdz	C		372 (M <sup>+</sup> +1)
N-a-11	NB1	Int.n-3	BRN6	Bn	H	Me	H	1Me-5-1HIdz	C		400 (M <sup>+</sup> +1)
N-a-12	NA	N-a-11		Bn	H	H	H	1Me-5-1HIdz	C		386 (M <sup>+</sup> +1)
N-a-13	NB1	Int.n-3	BRN7	Bn	H	Me	H	1Et-5-1HIdz	C		414 (M <sup>+</sup> +1)
N-a-14	NA	N-a-13		Bn	H	H	H	1Et-5-1HIdz	C		400 (M <sup>+</sup> +1)
N-a-15	NB1	Int.n-3	BRN8	Bn	H	Me	H	2Me-5-2HIdz	C		400 (M <sup>+</sup> +1)
N-a-16	NA	N-a-15		Bn	H	H	H	2Me-5-2HIdz	C		386 (M <sup>+</sup> +1)
N-a-17	NB1	Int.n-3	BRN9	Bn	H	Me	H	5-Bzt	C		403 (M <sup>+</sup> +1)
N-a-18	NA	N-a-17		Bn	H	H	H	5-Bzt	C		389 (M <sup>+</sup> +1)
N-a-19	NB1	Int.n-3	BRN10	Bn	H	Me	H	3-Qu	C		397 (M <sup>+</sup> +1)
N-a-20	NA	N-a-19		Bn	H	H	H	3-Qu	C		383 (M <sup>+</sup> +1)
N-a-21	NB1	Int.n-3	BRN11	Bn	H	Me	H	6-Qu	C		397 (M <sup>+</sup> +1)
N-a-22	NA	N-a-21		Bn	H	H	H	6-Qu	C		383 (M <sup>+</sup> +1)
N-a-23	NB1	Int.n-3	BRN12	Bn	H	Me	H	6-IQ	C		397 (M <sup>+</sup> +1)
N-a-24	NA	N-a-23		Bn	H	H	H	6-IQ	C		383 (M <sup>+</sup> +1)
N-a-25	NC1	N-a-1	CHO1	Bn	Me	Me	H	2-Nap	C		410 (M <sup>+</sup> +1)
N-a-26	NA	N-a-25		Bn	Me	H	H	2-Nap	C		396 (M <sup>+</sup> +1)
N-a-27	NC1	N-a-1	CHO2	Bn	Et	Me	H	2-Nap	C		424 (M <sup>+</sup> +1)
N-a-28	NA	N-a-27		Bn	Et	H	H	2-Nap	C		410 (M <sup>+</sup> +1)
N-a-29	NC1	N-a-3	CHO1	Bn	Me	Me	H	5-1Ind	C		399 (M <sup>+</sup> +1)
N-a-30	NA	N-a-29		Bn	Me	H	H	5-1Ind	C		384 (M <sup>+</sup> +1)
N-a-31	NC1	N-a-5	CHO1	Bn	Me	Me	H	1Me-5-Ind	C		413 (M <sup>+</sup> +1)
N-a-32	NA	N-a-31		Bn	Me	H	H	1Me-5-Ind	C		399 (M <sup>+</sup> +1)
N-a-33	NB1	Int.n-4	BRA1	4FBn	H	Me	H	2-Nap	C		414 (M <sup>+</sup> +1)
N-a-34	NA	N-a-33		4FBn	H	H	H	2-Nap	C		400 (M <sup>+</sup> +1)
N-a-35	NB1	Int.n-4	BRA2	4FBn	H	Me	H	5-1Ind	D	5.20	403 (M <sup>+</sup> +1)
N-a-36	NA	N-a-35		4FBn	H	H	H	5-1Ind	D	4.73	389 (M <sup>+</sup> +1)
N-a-37	NB1	Int.n-4	BRA3	4FBn	H	Me	H	1Me-5-Ind	D	5.51	417 (M <sup>+</sup> +1)
N-a-38	NA	N-a-37		4FBn	H	H	H	1Me-5-Ind	D	4.78	403 (M <sup>+</sup> +1)
N-a-39	NB1	Int.n-4	BRA5	4FBn	H	Me	H	5-1HIdz	D	4.60	404 (M <sup>+</sup> +1)
N-a-40	NA	N-a-39		4FBn	H	H	H	5-1HIdz	C		390 (M <sup>+</sup> +1)
N-a-41	NB1	Int.n-4	BRA6	4FBn	H	Me	H	1Me-5-1HIdz	A	4.85	418 (M <sup>+</sup> +1)
N-a-42	NA	N-a-41		4FBn	H	H	H	1Me-5-1HIdz	A	4.14	404 (M <sup>+</sup> +1)
N-a-43	NB1	Int.n-4	BRA10	4FBn	H	Me	H	3-Qu	D	4.72	415 (M <sup>+</sup> +1)
N-a-44	NA	N-a-43		4FBn	H	H	H	3-Qu	C		401 (M <sup>+</sup> +1)







Table-N-A-2

Exp.	Syn	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									method	RTime	Mass
N-a-45	NC2	N-a-35	CHO1	4FBn	Me	Me	H	5-Ind	D	4.17	417 (M <sup>+</sup> +1)
N-a-46	NA	N-a-45		4FBn	Me	H	H	5-Ind	D	3.38	403 (M <sup>+</sup> +1)
N-a-47	NC2	N-a-37	CHO1	4FBn	Me	Me	H	1Me-5-Ind	C		431 (M <sup>+</sup> +1)
N-a-48	NA	N-a-47		4FBn	Me	H	H	1Me-5-Ind	C		418 (M <sup>+</sup> +1)
N-a-49	NC2	N-a-41	CHO1	4FBn	Me	Me	H	1Me-5-1HIdz	C		432 (M <sup>+</sup> +1)
N-a-50	NA	N-a-49		4FBn	Me	H	H	1Me-5-1HIdz	C		418 (M <sup>+</sup> +1)
N-a-51	NC2	N-a-37	CHO2	4FBn	Et	Me	H	1Me-5-Ind	C		445 (M <sup>+</sup> +1)
N-a-52	NA	N-a-51		4FBn	Et	H	H	1Me-5-Ind	C		431 (M <sup>+</sup> +1)
N-a-53	NC2	N-a-39	CHO2	4FBn	Et	Me	H	5-1Idz	C		433 (M <sup>+</sup> +1)
N-a-54	NA	N-a-53		4FBn	Et	H	H	5-1Idz	C		419 (M <sup>+</sup> +1)
N-a-55	NB1	Int.n-5		2FBn	H	Me	H	2-Nap	C		414 (M <sup>+</sup> +1)
N-a-56	NA	N-a-55		2FBn	H	H	H	2-Nap	C		400 (M <sup>+</sup> +1)
N-a-57	NB1	Int.n-5		2FBn	H	Me	H	1Me-5-Ind	C		417 (M <sup>+</sup> +1)
N-a-58	NA	N-a-57		2FBn	H	H	H	1Me-5-Ind	C		403 (M <sup>+</sup> +1)
N-a-59	NB1	Int.n-5		2FBn	H	Me	H	1Me-5-1HIdz	C		418 (M <sup>+</sup> +1)
N-a-60	NA	N-a-59		2FBn	H	H	H	1Me-5-1HIdz	C		404 (M <sup>+</sup> +1)
N-a-61	NC2	N-a-59	CHO1	2FBn	Me	Me	H	1Me-5-1HIdz	C		432 (M <sup>+</sup> +1)
N-a-62	NA	N-a-61		2FBn	Me	H	H	1Me-5-1HIdz	C		418 (M <sup>+</sup> +1)
N-a-63	NB1	Int.n-6		3FBn	H	Me	H	2-Nap	C		414 (M <sup>+</sup> +1)
N-a-64	NA	N-a-63		3FBn	H	H	H	2-Nap	C		400 (M <sup>+</sup> +1)
N-a-65	NB1	Int.n-6		3FBn	H	Me	H	5-1Ind	C		403 (M <sup>+</sup> +1)
N-a-66	NA	N-a-65		3FBn	H	H	H	5-1Ind	C		389 (M <sup>+</sup> +1)
N-a-67	NB1	Int.n-6		3FBn	H	Me	H	1Me-5-Ind	C		417 (M <sup>+</sup> +1)
N-a-68	NA	N-a-67		3FBn	H	H	H	1Me-5-Ind	C		403 (M <sup>+</sup> +1)
N-a-69	NC2	N-a-67	CHO1	3FBn	Me	Me	H	1Me-5-Ind	C		431 (M <sup>+</sup> +1)
N-a-70	NA	N-a-69		3FBn	Me	H	H	1Me-5-Ind	C		417 (M <sup>+</sup> +1)
N-a-71	NC1	Int.n-7	CHO12	2ClBn	H	Me	H	2-Nap	C		430 (M <sup>+</sup> +1)
N-a-72	NA	N-a-71		2ClBn	H	H	H	2-Nap	C		416 (M <sup>+</sup> +1)
N-a-73	NC1	Int.n-7	CHO13	2BrBn	H	Me	H	2-Nap	C		475 (M <sup>+</sup> +1)
N-a-74	NA	N-a-73		2BrBn	H	H	H	2-Nap	C		461 (M <sup>+</sup> +1)
N-a-75	NC1	Int.n-7	CHO14	2,3DFBn	H	Me	H	2-Nap	C		432 (M <sup>+</sup> +1)
N-a-76	NA	N-a-75		2,3DFBn	H	H	H	2-Nap	C		418 (M <sup>+</sup> +1)
N-a-77	NC1	Int.n-7	CHO21	2-FR	H	Me	H	2-Nap	C		386 (M <sup>+</sup> +1)
N-a-78	NA	N-a-77		2-FR	H	H	H	2-Nap	C		372 (M <sup>+</sup> +1)
N-a-79	NC1	Int.n-7	CHO20	3-TF	H	Me	H	2-Nap	C		402 (M <sup>+</sup> +1)
N-a-80	NA	N-a-79		3-TF	H	H	H	2-Nap	C		388 (M <sup>+</sup> +1)
N-a-81	NC1	Int.n-7	CHO17	2CF3Bn	H	Me	H	2-Nap	C		464 (M <sup>+</sup> +1)
N-a-82	NA	N-a-80		2CF3Bn	H	H	H	2-Nap	C		450 (M <sup>+</sup> +1)
N-a-83	NC1	Int.n-8	CHO12	2ClBn	H	Me	H	5-1Ind	C		302 (M <sup>+</sup> +1)
N-a-84	NA	N-a-80		2ClBn	H	H	H	5-1Ind	C		288 (M <sup>+</sup> +1)
N-a-85	NC2	N-a-80	CHO1	2ClBn	Me	Me	H	5-1Ind	C		316 (M <sup>+</sup> +1)
N-a-86	NA	N-a-85		2ClBn	Me	H	H	5-1Ind	C		302 (M <sup>+</sup> +1)
N-a-87	NC1	Int.n-8	CHO14	2,3DFBn	H	Me	H	5-1Ind	C		304 (M <sup>+</sup> +1)
N-a-88	NA	N-a-87		2,3DFBn	H	H	H	5-1Ind	C		290 (M <sup>+</sup> +1)
N-a-89	NC1	Int.n-8	CHO16	4PhBn	H	Me	H	5-1Ind	C		344 (M <sup>+</sup> +1)
N-a-90	NA	N-a-89		4PhBn	H	H	H	5-1Ind	C		330 (M <sup>+</sup> +1)

Table-N-A-3

Exp.	Syn	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									method	RTime	Mass
N-a-91	NC1	Int.n-8	CHO19	2-TF	H	Me	H	5-Ind	C		391 (M <sup>+</sup> +1)
N-a-92	NA	N-a-91		2-TF	H	H	H	5-Ind	C		377 (M <sup>+</sup> +1)
N-a-93	NC1	Int.n-8	CHO17	2CF3Bn	H	Me	H	5-Ind	C		453 (M <sup>+</sup> +1)
N-a-94	NA	N-a-93		2CF3Bn	H	H	H	5-Ind	C		439 (M <sup>+</sup> +1)
N-a-95	NC1	Int.n-8	CHO18	2,3DCIBn	H	Me	H	5-Ind	C		454 (M <sup>+</sup> +1)
N-a-96	NA	N-a-71		2,3DCIBn	H	H	H	5-Ind	C		440 (M <sup>+</sup> +1)
N-a-97	NC1	Int.n-9	CHO13	2BrBn	H	Me	H	1Me-5-Ind	C		478 (M <sup>+</sup> +1)
N-a-98	NA	N-a-97		2BrBn	H	H	H	1Me-5-Ind	C		464 (M <sup>+</sup> +1)
N-a-99	NC1	Int.n-9	CHO15	3,4DFBn	H	Me	H	1Me-5-Ind	C		435 (M <sup>+</sup> +1)
N-a-100	NA	N-a-99		3,4DFBn	H	H	H	1Me-5-Ind	C		421 (M <sup>+</sup> +1)
N-a-101	NC1	Int.n-9	CHO16	4PhBn	H	Me	H	1Me-5-Ind	C		475 (M <sup>+</sup> +1)
N-a-102	NA	N-a-101		4PhBn	H	H	H	1Me-5-Ind	C		461 (M <sup>+</sup> +1)
N-a-103	NC1	Int.n-9	CHO21	2-FR	H	Me	H	1Me-5-Ind	C		389 (M <sup>+</sup> +1)
N-a-104	NA	N-a-103		2-FR	H	H	H	1Me-5-Ind	C		375 (M <sup>+</sup> +1)
N-a-105	NC1	Int.n-9	CHO20	3-TF	H	Me	H	1Me-5-Ind	C		405 (M <sup>+</sup> +1)
N-a-106	NA	N-a-105		3-TF	H	H	H	1Me-5-Ind	C		391 (M <sup>+</sup> +1)
N-a-107	NC1	Int.n-9	CHO18	2,3DCIBn	H	Me	H	1Me-5-Ind	C		468 (M <sup>+</sup> +1)
N-a-108	NA	N-a-107		2,3DCIBn	H	H	H	1Me-5-Ind	C		454 (M <sup>+</sup> +1)
N-a-109	NC1	Int.n-10	CHO13	2BrBn	H	Me	H	5-1HIdz	C		465 (M <sup>+</sup> +1)
N-a-110	NA	N-a-109		2BrBn	H	H	H	5-1HIdz	C		451 (M <sup>+</sup> +1)
N-a-111	NC1	Int.n-10	CHO15	3,4DFBn	H	Me	H	5-1HIdz	C		422 (M <sup>+</sup> +1)
N-a-112	NA	N-a-111		3,4DFBn	H	H	H	5-1HIdz	C		408 (M <sup>+</sup> +1)
N-a-113	NC2	N-a-111	CHO1	3,4DFBn	Me	Me	H	5-1HIdz	C		436 (M <sup>+</sup> +1)
N-a-114	NA	N-a-113		3,4DFBn	Me	H	H	5-1HIdz	C		422 (M <sup>+</sup> +1)
N-a-115	NC1	Int.n-10	CHO21	2-FR	H	Me	H	5-1HIdz	C		376 (M <sup>+</sup> +1)
N-a-116	NA	N-a-115		2-FR	H	H	H	5-1HIdz	C		362 (M <sup>+</sup> +1)
N-a-117	NC1	Int.n-10	CHO20	3-TF	H	Me	H	5-1HIdz	C		392 (M <sup>+</sup> +1)
N-a-118	NA	N-a-116		3-TF	H	H	H	5-1HIdz	C		378 (M <sup>+</sup> +1)
N-a-119	NC1	Int.n-10	CHO17	2CF3Bn	H	Me	H	5-1HIdz	C		454 (M <sup>+</sup> +1)
N-a-120	NA	N-a-120		2CF3Bn	H	H	H	5-1HIdz	C		440 (M <sup>+</sup> +1)
N-a-121	NC1	Int.n-10	CHO18	2,3DCIBn	H	Me	H	1Me-5-1HIdz	C		469 (M <sup>+</sup> +1)
N-a-122	NA	N-a-122		2,3DCIBn	H	H	H	1Me-5-1HIdz	C		455 (M <sup>+</sup> +1)
N-a-123	NC1	Int.n-11	CHO12	2CIBn	H	Me	H	1Me-5-1HIdz	C		434 (M <sup>+</sup> +1)
N-a-124	NA	N-a-123		2CIBn	H	H	H	1Me-5-1HIdz	C		420 (M <sup>+</sup> +1)
N-a-125	NC2	N-a-123	CHO1	2CIBn	Me	Me	H	1Me-5-1HIdz	C		448 (M <sup>+</sup> +1)
N-a-126	NA	N-a-125		2CIBn	Me	H	H	1Me-5-1HIdz	C		434 (M <sup>+</sup> +1)
N-a-127	NC1	Int.n-11	CHO14	2,3DFBn	H	Me	H	1Me-5-1HIdz	C		436 (M <sup>+</sup> +1)
N-a-128	NA	N-a-127		2,3DFBn	H	H	H	1Me-5-1HIdz	C		422 (M <sup>+</sup> +1)
N-a-129	NC1	Int.n-11	CHO15	3,4DFBn	H	Me	H	1Me-5-1HIdz	C		436 (M <sup>+</sup> +1)
N-a-130	NA	N-a-129		3,4DFBn	H	H	H	1Me-5-1HIdz	C		422 (M <sup>+</sup> +1)
N-a-131	NC1	Int.n-11	CHO16	4PhBn	H	Me	H	1Me-5-1HIdz	C		476 (M <sup>+</sup> +1)
N-a-132	NA	N-a-131		4PhBn	H	H	H	1Me-5-1HIdz	C		462 (M <sup>+</sup> +1)
N-a-133	NC1	Int.n-11	CHO19	2-TF	H	Me	H	1Me-5-1HIdz	C		406 (M <sup>+</sup> +1)
N-a-134	NA	N-a-133		2-TF	H	H	H	1Me-5-1HIdz	C		392 (M <sup>+</sup> +1)
N-a-135	NC1	Int.n-11	CHO17	2CF3Bn	H	Me	H	1Me-5-1HIdz	C		468 (M <sup>+</sup> +1)
N-a-136	NA	N-a-135		2CF3Bn	H	H	H	1Me-5-1HIdz	C		454 (M <sup>+</sup> +1)

Table-N-A-4

Exp.	Syn	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									metho	RTmin	Mass
N-a-137	NE1	Int.n-9			H	Me	H	1Me-5-Ind	C		413 (M <sup>+</sup> +1)
N-a-138	NA	N-a-137			H	H	H	1Me-5-Ind	C		399 (M <sup>+</sup> +1)
N-a-139	NE1	Int.n-11			H	Me	H	1Me-5-1HIdz	D	5.06	414 (M <sup>+</sup> +1)
N-a-140	NA	N-a-139			H	H	H	1Me-5-1HIdz	D	4.30	400 (M <sup>+</sup> +1)
N-a-141	NE2	Int.n-11	2(4FPh)EtOH	2(4FPh)Et	H	Me	H	1Me-5-1HIdz	D	5.08	432 (M <sup>+</sup> +1)
N-a-142	NA	N-a-141		2(4FPh)Et	H	H	H	1Me-5-1HIdz	D	4.25	418 (M <sup>+</sup> +1)
N-a-143	NF	N-a-5	AcCl	Bn	Ac	Me	H	1Me-5-1HIdz	C		444 (M <sup>+</sup> +1)
N-a-144	NA	N-a-143		Bn	Ac	H	H	1Me-5-1HIdz	C		430 (M <sup>+</sup> +1)
N-a-145	NF	N-a-5	PhCOCl	Bn	PhC(O)	Me	H	1Me-5-Ind	C		504 (M <sup>+</sup> +1)
N-a-146	NA	N-a-145		Bn	PhC(O)	H	H	1Me-5-Ind	C		490 (M <sup>+</sup> +1)
N-a-147	NF	N-a-5	MeOCH <sub>2</sub> COCl	Bn	MeOCH <sub>2</sub> C(O)	Me	H	1Me-5-Ind	C		472 (M <sup>+</sup> +1)
N-a-148	NA	N-a-147		Bn	MeOCH <sub>2</sub> C(O)	H	H	1Me-5-Ind	C		458 (M <sup>+</sup> +1)
N-a-149	NF	N-a-5	MeOCOC(=O)Cl	Bn	MeOC(O)	Me	H	1Me-5-Ind	C		458 (M <sup>+</sup> +1)
N-a-150	NA	N-a-149		Bn	MeOC(O)	H	H	1Me-5-Ind	C		444 (M <sup>+</sup> +1)
N-a-151	NF	N-a-5	PhOCOC(=O)Cl	Bn	PhOC(O)	Me	H	1Me-5-Ind	C		520 (M <sup>+</sup> +1)
N-a-152	NA	N-a-151		Bn	PhOC(O)	H	H	1Me-5-Ind	C		506 (M <sup>+</sup> +1)
N-a-153	NF	N-a-5	NMe <sub>2</sub> COCl	Bn	Me <sub>2</sub> NC(O)	Me	H	1Me-5-Ind	C		471 (M <sup>+</sup> +1)
N-a-154	NA	N-a-153		Bn	Me <sub>2</sub> NC(O)	H	H	1Me-5-Ind	C		457 (M <sup>+</sup> +1)
N-a-155	NF	N-a-11	AcCl	Bn	Ac	Me	H	1Me-5-Ind	C		442 (M <sup>+</sup> +1)
N-a-156	NA	N-a-155		Bn	Ac	H	H	1Me-5-Ind	C		428 (M <sup>+</sup> +1)
N-a-157	NF	N-a-5	AcCl	4FBn	Ac	Me	H	1Me-5-Ind	C		461 (M <sup>+</sup> +1)
N-a-158	NA	N-a-157		4FBn	Ac	H	H	1Me-5-Ind	C		447 (M <sup>+</sup> +1)
N-a-159	NF	N-a-5	MeOCH <sub>2</sub> COCl	4FBn	MeOCH <sub>2</sub> C(O)	Me	H	1Me-5-Ind	C		491 (M <sup>+</sup> +1)
N-a-160	NA	N-a-159		4FBn	MeOCH <sub>2</sub> C(O)	H	H	1Me-5-Ind	C		477 (M <sup>+</sup> +1)
N-a-161	NF	N-a-5	MeOCOC(=O)Cl	4FBn	MeOC(O)	Me	H	1Me-5-Ind	C		477 (M <sup>+</sup> +1)
N-a-162	NA	N-a-161		4FBn	MeOC(O)	H	H	1Me-5-Ind	C		463 (M <sup>+</sup> +1)
N-a-163	NF	N-a-11	AcCl	4FBn	Ac	Me	H	1Me-5-1HIdz	C		462 (M <sup>+</sup> +1)
N-a-164	NA	N-a-163		4FBn	Ac	H	H	1Me-5-1HIdz	C		448 (M <sup>+</sup> +1)
N-a-165	NF	N-a-11	MeOCOC(=O)Cl	4FBn	MeOC(O)	Me	H	1Me-5-1HIdz	C		478 (M <sup>+</sup> +1)
N-a-166	NA	N-a-165		4FBn	MeOC(O)	H	H	1Me-5-1HIdz	C		464 (M <sup>+</sup> +1)

[Example N-b-1]

Synthesis of methyl 3-[4-(N-methylamino)-3-(naphthalen-2-yl)phenyl]propionate  
(Compound No. N-b-1) (Synthesis method ND1)

According to the procedure described in the synthesis method of Intermediate Int. n-7 (Synthesis method ND1) provided that the reaction was carried out for 2 hours, the compound of Example N-a-25 (100.3 mg) and 10% palladium/carbon (10.2 mg) were reacted and treated to obtain the title compound (Compound No. N-b-1, 89.7 mg).

[Example N-b-35]

Synthesis of methyl 3-[4-(N-ethylamino)-3-(naphthalen-2-yl)phenyl]propionate  
(Compound No. N-b-35) (Synthesis method NB1)

According to the procedure described in the synthesis method of the compound of Example N-a-1 (Synthesis method NB1) provided that the reaction was carried out for 17 hours, Intermediate n-12 (99.87 mg), 2-naphthaleneboronic acid (87.3 mg), 2 M aqueous sodium carbonate (350  $\mu$ l) and (Ph<sub>3</sub>P)<sub>4</sub>Pd (59.6 mg) were reacted and treated to obtain the title compound (Compound No. N-b-35, 103.5 mg).

[Example N-b-79]

Synthesis of methyl 3-[4-(N-n-butylamino)-3-(naphthalen-2-yl)phenyl]propionate  
(Compound No. N-b-79) (Synthesis method NC2)

According to the procedure described in the synthesis method of Intermediate n-3 provided that the reaction was carried out for 13 hours, Intermediate n-7 (164.7 mg) and n-butylaldehyde (38.5  $\mu$ l, KANTO), sodium triacetoxyborohydride (138.6 mg) and acetic acid (75  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. N-b-79, 161.3 mg).

[Example N-b-183]

Synthesis of methyl 3-[4-(N-acetyl-N-methylamino)-3-(naphthalen-2-yl)phenyl]propionate (Compound No. N-b-183) (Synthesis method NF)

According to the procedure described in the synthesis method of the compound of Example N-a-143 provided that the reaction was carried out for 18 hours, the compound of Example N-b-1 (22.7 mg), pyridine (23.8  $\mu$ l) and acetyl chloride (40  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. N-b-183, 16.3 mg).

[Example N-b-197]

Synthesis of 3-[4-(N-benzoyl-N-methylamino)-3-(naphthalen-2-yl)phenyl]propionic acid (Compound No. N-b-197) (Synthesis method NF)



According to the procedure described in the synthesis method of the compound of Example N-a-143 provided that the reaction was carried out for 14 hours, the compound of Example N-b-1 (21.8 mg), pyridine (23.8  $\mu$  l) and benzoyl chloride (345  $\mu$  l) were reacted and treated. A solution of the obtained residue in methanol (3 ml) was added with 2 N aqueous sodium hydroxide (100  $\mu$  l), and stirred at 60°C for 2 hours. The reaction mixture was concentrated under reduced pressure, then made acidic with 5% aqueous hydrochloric acid under ice cooling, and extracted with dichloromethane (5 ml). The organic layer was washed successively with saturated brine, and dried, and then the solvent was evaporated under reduced pressure to obtain the title compound (Compound No. N-b-197, 13.5 mg).

[Examples N-b-1 to N-b-212]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification including the examples described above are shown in Table-N-B-1 to Table-N-B-5. In the tables, the compound numbers are mentioned in the columns indicated as "Exp.". In the tables, used methods among the aforementioned synthesis methods are shown in the columns of "Syn" with symbols, the starting compounds 1 are mentioned in the columns of "SM1", and the starting compounds 2 are mentioned in the columns of "SM2".

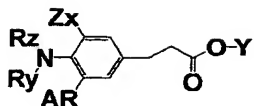


Table-N-B-1

Exp.	Syn	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									method	RTime	Mass
N-b-1	ND1	N-a-25		Me	H	Me	H	2-Nap	C		320 (M <sup>+</sup> +1)
N-b-2	NA	N-b-1		Me	H	H	H	2-Nap	C		306 (M <sup>+</sup> +1)
N-b-3	ND1	N-a-29		Me	H	Me	H	5-Ind	C		309 (M <sup>+</sup> +1)
N-b-4	NA	N-b-3		Me	H	H	H	5-Ind	C		295 (M <sup>+</sup> +1)
N-b-5	ND1	N-a-31		Me	H	Me	H	1Me-5-Ind	C		323 (M <sup>+</sup> +1)
N-b-6	NA	N-b-5		Me	H	H	H	1Me-5-Ind	C		309 (M <sup>+</sup> +1)
N-b-7	ND1	N-a-69		Me	H	Me	H	5-1HIdz	C		310 (M <sup>+</sup> +1)
N-b-8	NA	N-b-7		Me	H	H	H	5-1HIdz	C		296 (M <sup>+</sup> +1)
N-b-9	ND1	N-a-49		Me	H	Me	H	1Me-5-1HIdz	C		324 (M <sup>+</sup> +1)
N-b-10	NA	N-b-9		Me	H	H	H	1Me-5-1HIdz	C		310 (M <sup>+</sup> +1)
N-b-11	NC2	N-b-1	CHO1	Me	Me	Me	H	2-Nap	C		334 (M <sup>+</sup> +1)
N-b-12	NA	N-b-11		Me	Me	H	H	2-Nap	C		320 (M <sup>+</sup> +1)
N-b-13	NC2	N-b-1	CHO2	Me	Et	Me	H	2-Nap	C		348 (M <sup>+</sup> +1)
N-b-14	NA	N-b-13		Me	Et	H	H	2-Nap	C		334 (M <sup>+</sup> +1)
N-b-15	NC2	N-b-3	CHO1	Me	Me	Me	H	5-Ind	C		323 (M <sup>+</sup> +1)
N-b-16	NA	N-b-15		Me	Me	H	H	5-Ind	C		309 (M <sup>+</sup> +1)
N-b-17	NC2	N-b-5	CHO1	Me	Me	H	H	1Me-5-Ind	C		337 (M <sup>+</sup> +1)
N-b-18	NA	N-b-17		Me	Me	H	H	1Me-5-Ind	C		323 (M <sup>+</sup> +1)
N-b-19	NC2	N-b-9	CHO1	Me	Me	Me	H	1Me-5-1HIdz	C		338 (M <sup>+</sup> +1)
N-b-20	NA	N-b-19		Me	Me	H	H	1Me-5-1HIdz	C		324 (M <sup>+</sup> +1)
N-b-21	NB1	Int.n-12	BRA1	Et	H	Me	H	2-Nap	C		334 (M <sup>+</sup> +1)
N-b-22	NA	N-b-21		Et	H	H	H	2-Nap	C		320 (M <sup>+</sup> +1)
N-b-23	NB1	Int.n-12	BRA2	Et	H	Me	H	5-Ind	C		323 (M <sup>+</sup> +1)
N-b-24	NA	N-b-23		Et	H	H	H	5-Ind	C		309 (M <sup>+</sup> +1)
N-b-25	NB1	Int.n-12	BRA3	Et	H	Me	H	1Me-5-Ind	C		337 (M <sup>+</sup> +1)
N-b-26	NA	N-b-25		Et	H	H	H	1Me-5-Ind	C		323 (M <sup>+</sup> +1)
N-b-27	NB1	Int.n-12	BRA4	Et	H	Me	H	1Et-5-Ind	C		351 (M <sup>+</sup> +1)
N-b-28	NA	N-b-27		Et	H	H	H	1Et-5-Ind	C		337 (M <sup>+</sup> +1)
N-b-29	NB1	Int.n-12	BRA5	Et	H	Me	H	5-1HIdz	C		324 (M <sup>+</sup> +1)
N-b-30	NA	N-b-29		Et	H	H	H	5-1HIdz	C		310 (M <sup>+</sup> +1)
N-b-31	NB1	Int.n-12	BRA6	Et	H	Me	H	1Me-5-1HIdz	C		338 (M <sup>+</sup> +1)
N-b-32	NA	N-b-31		Et	H	H	H	1Me-5-1HIdz	C		324 (M <sup>+</sup> +1)
N-b-33	NB1	Int.n-12	BRA7	Et	H	Me	H	1Et-5-Idz	C		352 (M <sup>+</sup> +1)
N-b-34	NA	N-b-33		Et	H	H	H	1Et-5-Idz	C		338 (M <sup>+</sup> +1)
N-b-35	NB1	Int.n-12	BRA8	Et	H	Me	H	2Me-5-Idz	C		338 (M <sup>+</sup> +1)
N-b-36	NA	N-b-35		Et	H	H	H	2Me-5-Idz	C		324 (M <sup>+</sup> +1)
N-b-37	NB1	Int.n-12	BRA9	Et	H	Me	H	5-Bzt	C		341 (M <sup>+</sup> +1)
N-b-38	NA	N-b-37		Et	H	H	H	5-Bzt	C		327 (M <sup>+</sup> +1)
N-b-39	NB1	Int.n-12	BRA10	Et	H	Me	H	3-Qu	C		335 (M <sup>+</sup> +1)
N-b-40	NA	N-b-39		Et	H	H	H	3-Qu	C		321 (M <sup>+</sup> +1)
N-b-41	NB1	Int.n-12	BRA11	Et	H	Me	H	6-Qu	C		335 (M <sup>+</sup> +1)
N-b-42	NA	N-b-41		Et	H	H	H	6-Qu	C		321 (M <sup>+</sup> +1)
N-b-43	NC2	N-b-21	CHO2	Et	Et	Me	H	2-Nap	C		362 (M <sup>+</sup> +1)
N-b-44	NA	N-b-43		Et	Et	H	H	2-Nap	C		348 (M <sup>+</sup> +1)

Table-N-B-2

Exp.	Syn.	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									method	RTime	Mass
N-b-45	NC2	N-b-25	CHO2	Et	Et	Me	H	1Me-5-Ind	C		365 (M <sup>+</sup> +1)
N-b-46	NA	N-b-45		Et	Et	H	H	1Me-5-Ind	C		351 (M <sup>+</sup> +1)
N-b-47	NB1	Int.n-13		nPr	H	Me	H	5-Ind	C		337 (M <sup>+</sup> +1)
N-b-48	NA	N-b-47		nPr	H	H	H	5-Ind	C		323 (M <sup>+</sup> +1)
N-b-49	NB1	Int.n-13		nPr	H	Me	H	1Me-5-Ind	C		351 (M <sup>+</sup> +1)
N-b-50	NA	N-b-49		nPr	H	H	H	1Me-5-Ind	C		337 (M <sup>+</sup> +1)
N-b-51	NB1	Int.n-13		nPr	H	Me	H	5-1HIdz	C		338 (M <sup>+</sup> +1)
N-b-52	NA	N-b-51		nPr	H	H	H	5-1HIdz	C		324 (M <sup>+</sup> +1)
N-b-53	NB1	Int.n-13		nPr	H	Me	H	1Me-5-1HIdz	C		352 (M <sup>+</sup> +1)
N-b-54	NA	N-b-53		nPr	H	H	H	1Me-5-1HIdz	C		338 (M <sup>+</sup> +1)
N-b-55	NC2	N-b-47	CHO1	nPr	Me	Me	H	5-Ind	C		351 (M <sup>+</sup> +1)
N-b-56	NA	N-b-55		nPr	Me	H	H	5-Ind	C		337 (M <sup>+</sup> +1)
N-b-57	NC2	N-b-49	CHO1	nPr	Me	Me	H	1Me-5-Ind	C		365 (M <sup>+</sup> +1)
N-b-58	NA	N-b-57		nPr	Me	H	H	1Me-5-Ind	C		351 (M <sup>+</sup> +1)
N-b-59	NC2	N-b-51	CHO1	nPr	Me	Me	H	5-1HIdz	C		352 (M <sup>+</sup> +1)
N-b-60	NA	N-b-59		nPr	Me	H	H	5-1HIdz	C		338 (M <sup>+</sup> +1)
N-b-61	NC2	N-b-53	CHO1	nPr	Me	Me	H	1Me-5-1HIdz	C		366 (M <sup>+</sup> +1)
N-b-62	NA	N-b-61		nPr	Me	H	H	1Me-5-1HIdz	C		352 (M <sup>+</sup> +1)
N-b-63	NB1	Int.n-14	BRA2	iPr	H	Me	H	5-Ind	C		337 (M <sup>+</sup> +1)
N-b-64	NA	N-b-63		iPr	H	H	H	5-Ind	C		323 (M <sup>+</sup> +1)
N-b-65	NB1	Int.n-14	BRA3	iPr	H	Me	H	1Me-5-Ind	C		351 (M <sup>+</sup> +1)
N-b-66	NA	N-b-65		iPr	H	H	H	1Me-5-Ind	C		337 (M <sup>+</sup> +1)
N-b-67	NB1	Int.n-14	BRA5	iPr	H	Me	H	5-1HIdz	C		338 (M <sup>+</sup> +1)
N-b-68	NA	N-b-67		iPr	H	H	H	5-1HIdz	C		324 (M <sup>+</sup> +1)
N-b-69	NB1	Int.n-14	BRA6	iPr	H	Me	H	1Me-5-1HIdz	C		352 (M <sup>+</sup> +1)
N-b-70	NA	N-b-69		iPr	H	H	H	1Me-5-1HIdz	C		338 (M <sup>+</sup> +1)
N-b-71	NC2	N-b-63	CHO1	iPr	Me	Me	H	5-Ind	C		351 (M <sup>+</sup> +1)
N-b-72	NA	N-b-71		iPr	Me	H	H	5-Ind	C		337 (M <sup>+</sup> +1)
N-b-73	NC2	N-b-65	CHO1	iPr	Me	Me	H	1Me-5-Ind	C		365 (M <sup>+</sup> +1)
N-b-74	NA	N-b-73		iPr	Me	H	H	1Me-5-Ind	C		351 (M <sup>+</sup> +1)
N-b-75	NC1	N-b-67	CHO1	iPr	Me	Me	H	5-1HIdz	C		352 (M <sup>+</sup> +1)
N-b-76	NA	N-b-75		iPr	Me	H	H	5-1HIdz	C		338 (M <sup>+</sup> +1)
N-b-77	NC1	N-b-69	CHO1	iPr	Me	Me	H	1Me-5-1HIdz	C		366 (M <sup>+</sup> +1)
N-b-78	NA	N-b-77		iPr	Me	H	H	1Me-5-1HIdz	C		352 (M <sup>+</sup> +1)
N-b-79	NB1	Int.n-7	BRA1	nBu	H	Me	H	2-Nap	C		362 (M <sup>+</sup> +1)
N-b-80	NA	N-b-79		nBu	H	H	H	2-Nap	C		348 (M <sup>+</sup> +1)
N-b-81	NB1	Int.n-8	BRA2	nBu	H	Me	H	5-Ind	C		351 (M <sup>+</sup> +1)
N-b-82	NA	N-b-81		nBu	H	H	H	5-Ind	C		337 (M <sup>+</sup> +1)
N-b-83	NB1	Int.n-10	BRA5	nBu	H	Me	H	5-1HIdz	C		352 (M <sup>+</sup> +1)
N-b-84	NA	N-b-83		nBu	H	H	H	5-1HIdz	C		338 (M <sup>+</sup> +1)
N-b-85	NB1	Int.n-11	BRA6	nBu	H	Me	H	1Me-5-1HIdz	C		366 (M <sup>+</sup> +1)
N-b-86	NA	N-b-85		nBu	H	H	H	1Me-5-1HIdz	C		352 (M <sup>+</sup> +1)
N-b-87	NC1	N-b-79	CHO1	nBu	Me	Me	H	2-Nap	C		376 (M <sup>+</sup> +1)
N-b-88	NA	N-b-87		nBu	Me	H	H	2-Nap	C		351 (M <sup>+</sup> +1)
N-b-89	NC1	N-b-81	CHO1	nBu	Me	Me	H	5-Ind	C		365 (M <sup>+</sup> +1)
N-b-90	NA	N-b-89		nBu	Me	H	H	5-Ind	C		351 (M <sup>+</sup> +1)









Table-N-B-3

Exp.	Syn	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									method	RTime	Mass
N-b-91	NC1	N-b-83	CHO1	nBu	Me	Me	H	5-1HIdz	C		366 (M <sup>+</sup> +1)
N-b-92	NA	N-b-91		nBu	Me	H	H	5-1HIdz	C		352 (M <sup>+</sup> +1)
N-b-93	NC1	N-b-85	CHO1	nBu	Me	Me	H	1Me-5-1HIdz	C		380 (M <sup>+</sup> +1)
N-b-94	NA	N-b-93		nBu	Me	H	H	1Me-5-1HIdz	C		366 (M <sup>+</sup> +1)
N-b-95	NC2	N-b-81	CHO2	nBu	Et	Me	H	5-Ind	C		379 (M <sup>+</sup> +1)
N-b-96	NA	N-b-95		nBu	Et	H	H	5-Ind	C		365 (M <sup>+</sup> +1)
N-b-97	NC2	N-b-85	CHO2	nBu	Et	Me	H	1Me-5-1HIdz	C		394 (M <sup>+</sup> +1)
N-b-98	NA	N-b-97		nBu	Et	H	H	1Me-5-1HIdz	C		380 (M <sup>+</sup> +1)
N-b-99	NC2	Int.n-9	CHO7	iBu	H	Me	H	1Me-5-Ind	C		365 (M <sup>+</sup> +1)
N-b-100	NA	N-b-99		iBu	H	H	H	1Me-5-Ind	C		351 (M <sup>+</sup> +1)
N-b-101	NC2	Int.n-10	CHO7	iBu	H	Me	H	5-1HIdz	C		352 (M <sup>+</sup> +1)
N-b-102	NA	N-b-101		iBu	H	H	H	5-1HIdz	C		338 (M <sup>+</sup> +1)
N-b-103	NC2	Int.n-11	CHO7	iBu	H	Me	H	1Me-5-1HIdz	C		366 (M <sup>+</sup> +1)
N-b-104	NA	N-b-103		iBu	H	H	H	1Me-5-1HIdz	C		352 (M <sup>+</sup> +1)
N-b-105	NC2	Int.n-15	BRA11	iBu	H	Me	H	6-Qu	C		363 (M <sup>+</sup> +1)
N-b-106	NA	N-b-105		iBu	H	H	H	6-Qu	C		349 (M <sup>+</sup> +1)
N-b-107	NC1	N-b-99	CHO1	iBu	Me	Me	H	1Me-5-Ind	C		379 (M <sup>+</sup> +1)
N-b-108	NA	N-b-107		iBu	Me	H	H	1Me-5-Ind	C		365 (M <sup>+</sup> +1)
N-b-109	NC1	N-b-103	CHO1	iBu	Me	Me	H	1Me-5-1HIdz	C		380 (M <sup>+</sup> +1)
N-b-110	NA	N-b-109		iBu	Me	H	H	1Me-5-1HIdz	C		366 (M <sup>+</sup> +1)
N-b-111	NC1	N-b-105	CHO1	iBu	Me	Me	H	6-Qu	C		377 (M <sup>+</sup> +1)
N-b-112	NA	N-b-111		iBu	Me	H	H	6-Qu	C		363 (M <sup>+</sup> +1)
N-b-113	NC2	N-b-99	CHO2	iBu	Et	Me	H	1Me-5-Ind	C		393 (M <sup>+</sup> +1)
N-b-114	NA	N-b-113		iBu	Et	H	H	1Me-5-Ind	C		379 (M <sup>+</sup> +1)
N-b-115	NC2	N-b-101	CHO2	iBu	Et	Me	H	5-1HIdz	C		380 (M <sup>+</sup> +1)
N-b-116	NA	N-b-115		iBu	Et	H	H	5-1HIdz	C		366 (M <sup>+</sup> +1)
N-b-117	NC2	N-b-103	CHO2	iBu	Et	Me	H	1Me-5-1HIdz	C		394 (M <sup>+</sup> +1)
N-b-118	NA	N-b-117		iBu	Et	H	H	1Me-5-1HIdz	C		380 (M <sup>+</sup> +1)
N-b-119	NB1	Int.n-16	BRA1	cPen	H	Me	H	2-Nap	C		374 (M <sup>+</sup> +1)
N-b-120	NA	N-b-119		cPen	H	H	H	2-Nap	C		360 (M <sup>+</sup> +1)
N-b-121	NB1	Int.n-16	BRA2	cPen	H	Me	H	5-Ind	C		363 (M <sup>+</sup> +1)
N-b-122	NA	N-b-121		cPen	H	H	H	5-Ind	C		349 (M <sup>+</sup> +1)
N-b-123	NB1	Int.n-9	BRA3	cPen	H	Me	H	1Me-5-Ind	C		377 (M <sup>+</sup> +1)
N-b-124	NA	N-b-123		cPen	H	H	H	1Me-5-Ind	C		363 (M <sup>+</sup> +1)
N-b-125	NB1	Int.n-16	BRA5	cPen	H	Me	H	5-1HIdz	C		364 (M <sup>+</sup> +1)
N-b-126	NA	N-b-125		cPen	H	H	H	5-1HIdz	C		350 (M <sup>+</sup> +1)
N-b-127	NB1	Int.n-11	BRA6	cPen	H	Me	H	1Me-5-1HIdz	C		378 (M <sup>+</sup> +1)
N-b-128	NA	N-b-127		cPen	H	H	H	1Me-5-1HIdz	C		364 (M <sup>+</sup> +1)
N-b-129	NB1	Int.n-16	BRA11	cPen	H	Me	H	6-Qu	C		375 (M <sup>+</sup> +1)
N-b-130	NA	N-b-129		cPen	H	H	H	6-Qu	C		361 (M <sup>+</sup> +1)
N-b-131	NB1	Int.n-16	BRA9	cPen	H	Me	H	5-Bzt	C		381 (M <sup>+</sup> +1)
N-b-132	NA	N-b-131		cPen	H	H	H	5-Bzt	C		367 (M <sup>+</sup> +1)
N-b-133	NC1	N-b-121	CHO1	cPen	Me	Me	H	5-Ind	C		377 (M <sup>+</sup> +1)
N-b-134	NA	N-b-133		cPen	Me	H	H	5-Ind	C		363 (M <sup>+</sup> +1)
N-b-135	NC1	N-b-123	CHO1	cPen	Me	Me	H	1Me-5-Ind	C		391 (M <sup>+</sup> +1)
N-b-136	NA	N-b-135		cPen	Me	H	H	1Me-5-Ind	C		377 (M <sup>+</sup> +1)

Table-N-B-4

Exp.	Syn.	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									method	RTime	Mass
N-b-137	NC1	N-b-127	CHO1	cPen	Me	Me	H	1Me-5-1HIdz	C		392 (M <sup>+</sup> +1)
N-b-138	NA	N-b-137		cPen	Me	H	H	1Me-5-1HIdz	C		378 (M <sup>+</sup> +1)
N-b-139	NC2	N-b-123	CHO2	cPen	Et	Me	H	1Me-5-Ind	C		405 (M <sup>+</sup> +1)
N-b-140	NA	N-b-139		cPen	Et	H	H	1Me-5-Ind	C		391 (M <sup>+</sup> +1)
N-b-141	NC2	N-b-131	CHO2	cPen	Et	Me	H	5-Bzt	C		409 (M <sup>+</sup> +1)
N-b-142	NA	N-b-141		cPen	Et	H	H	5-Bzt	C		395 (M <sup>+</sup> +1)
N-b-143	NB1	Int.n-17	BRA1	cHex	H	Me	H	2-Nap	C		388 (M <sup>+</sup> +1)
N-b-144	NA	N-b-143		cHex	H	H	H	2-Nap	C		374 (M <sup>+</sup> +1)
N-b-145	NB1	Int.n-17	BRA2	cHex	H	Me	H	5-Ind	C		377 (M <sup>+</sup> +1)
N-b-146	NA	N-b-145		cHex	H	H	H	5-Ind	C		363 (M <sup>+</sup> +1)
N-b-147	NB1	Int.n-9	BRA3	cHex	H	Me	H	1Me-5-Ind	C		391 (M <sup>+</sup> +1)
N-b-148	NA	N-b-147		cHex	H	H	H	1Me-5-Ind	C		377 (M <sup>+</sup> +1)
N-b-149	NB1	Int.n-17	BRA5	cHex	H	Me	H	5-1HIdz	C		378 (M <sup>+</sup> +1)
N-b-150	NA	N-b-149		cHex	H	H	H	5-1HIdz	C		364 (M <sup>+</sup> +1)
N-b-151	NB1	Int.n-17	BRA6	cHex	H	Me	H	1Me-5-1HIdz	C		392 (M <sup>+</sup> +1)
N-b-152	NA	N-b-151		cHex	H	H	H	1Me-5-1HIdz	C		378 (M <sup>+</sup> +1)
N-b-153	NB1	Int.n-17	BRA10	cHex	H	Me	H	3-Qu	C		389 (M <sup>+</sup> +1)
N-b-154	NA	N-b-153		cHex	H	H	H	3-Qu	C		375 (M <sup>+</sup> +1)
N-b-155	NC1	N-b-143	CHO1	cHex	Me	Me	H	2-Nap	C		402 (M <sup>+</sup> +1)
N-b-156	NA	N-b-155		cHex	Me	H	H	2-Nap	C		388 (M <sup>+</sup> +1)
N-b-157	NC1	N-b-147	CHO1	cHex	Me	Me	H	1Me-5-Ind	C		405 (M <sup>+</sup> +1)
N-b-158	NA	N-b-157		cHex	Me	H	H	1Me-5-Ind	C		391 (M <sup>+</sup> +1)
N-b-159	NC1	N-b-149	CHO1	cHex	Me	Me	H	5-1HIdz	C		392 (M <sup>+</sup> +1)
N-b-160	NA	N-b-159		cHex	Me	H	H	5-1HIdz	C		378 (M <sup>+</sup> +1)
N-b-161	NC1	N-b-151	CHO1	cHex	Me	Me	H	1Me-5-1HIdz	C		406 (M <sup>+</sup> +1)
N-b-162	NA	N-b-161		cHex	Me	H	H	1Me-5-1HIdz	C		392 (M <sup>+</sup> +1)
N-b-163	NC2	N-b-143	CHO2	cHex	Et	Me	H	2-Nap	C		416 (M <sup>+</sup> +1)
N-b-164	NA	N-b-163		cHex	Et	H	H	2-Nap	C		402 (M <sup>+</sup> +1)
N-b-165	NC2	N-b-153	CHO2	cHex	Et	Me	H	3-Qu	C		417 (M <sup>+</sup> +1)
N-b-166	NA	N-b-165		cHex	Et	H	H	3-Qu	C		403 (M <sup>+</sup> +1)
N-b-167	NB1	Int.n-18	BRA2	2(Me)cHex	H	Me	H	5-Ind	C		391 (M <sup>+</sup> +1)
N-b-168	NA	N-b-167		2(Me)cHex	H	H	H	5-Ind	C		377 (M <sup>+</sup> +1)
N-b-169	NB1	Int.n-18	BRA3	2(Me)cHex	H	Me	H	1Me-5-Ind	C		405 (M <sup>+</sup> +1)
N-b-170	NA	N-b-169		2(Me)cHex	H	H	H	1Me-5-Ind	C		391 (M <sup>+</sup> +1)
N-b-171	NB1	Int.n-18	BRA5	2(Me)cHex	H	Me	H	5-1HIdz	C		392 (M <sup>+</sup> +1)
N-b-172	NA	N-b-171		2(Me)cHex	H	H	H	5-1HIdz	C		378 (M <sup>+</sup> +1)
N-b-173	NB1	Int.n-18	BRA6	2(Me)cHex	H	Me	H	1Me-5-1HIdz	C		406 (M <sup>+</sup> +1)
N-b-174	NA	N-b-173		2(Me)cHex	H	H	H	1Me-5-1HIdz	C		392 (M <sup>+</sup> +1)
N-b-175	NC2	Int.n-8	CHO25	2-Indane	H	Me	H	5-Ind	C		411 (M <sup>+</sup> +1)
N-b-176	NA	N-b-175		2-Indane	H	H	H	5-Ind	C		397 (M <sup>+</sup> +1)
N-b-177	NC2	Int.n-9	CHO25	2-Indane	H	Me	H	1Me-5-Ind	C		425 (M <sup>+</sup> +1)
N-b-178	NA	N-b-177		2-Indane	H	H	H	1Me-5-Ind	C		411 (M <sup>+</sup> +1)
N-b-179	NC2	Int.n-10	CHO25	2-Indane	H	Me	H	5-1HIdz	C		412 (M <sup>+</sup> +1)
N-b-180	NA	N-b-179		2-Indane	H	H	H	5-1HIdz	C		398 (M <sup>+</sup> +1)
N-b-181	NC2	Int.n-11	CHO25	2-Indane	H	Me	H	1Me-5-1HIdz	C		426 (M <sup>+</sup> +1)
N-b-182	NA	N-b-181		2-Indane	H	H	H	1Me-5-1HIdz	C		412 (M <sup>+</sup> +1)

Table-N-B-5

Exp.	Syn	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									method	RTTime	Mass
N-b-183	NF	N-b-1	AcCl	Me	Ac	Me	H	2-Nap	C		364 (M <sup>+</sup> +1)
N-b-184	NA	N-b-183		Me	Ac	H	H	2-Nap	C		350 (M <sup>+</sup> +1)
N-b-185	NF	N-b-5	AcCl	Me	Ac	Me	H	1Me-5-Ind	C		367 (M <sup>+</sup> +1)
N-b-186	NA	N-b-185		Me	Ac	H	H	1Me-5-Ind	C		353 (M <sup>+</sup> +1)
N-b-187	NF	N-b-11	AcCl	Me	Ac	Me	H	1Me-5-1Hldz	C		368 (M <sup>+</sup> +1)
N-b-188	NA	N-b-187		Me	Ac	H	H	1Me-5-1Hldz	C		354 (M <sup>+</sup> +1)
N-b-189	NF	Intn-11	AcCl	Ac	Ac	Me	H	1Me-5-1Hldz	C		394 (M <sup>+</sup> +1)
N-b-190	NA	N-b-189		Ac	Ac	H	H	1Me-5-1Hldz	C		380 (M <sup>+</sup> +1)
N-b-191	NF	Intn-18	MeOCOCi	Me	MeOC(O)	Me	H	2-Nap	C		380 (M <sup>+</sup> +1)
N-b-192	NA	N-b-167		Me	MeOC(O)	H	H	2-Nap	C		366 (M <sup>+</sup> +1)
N-b-193	NF	Intn-18	MeOCOCi	Me	MeOC(O)	Me	H	1Me-5-Ind	C		383 (M <sup>+</sup> +1)
N-b-194	NA	N-b-169		Me	MeOC(O)	H	H	1Me-5-Ind	C		369 (M <sup>+</sup> +1)
N-b-195	NF	Intn-18	MeOCOCi	Me	MeOC(O)	Me	H	1Me-5-1Hldz	C		384 (M <sup>+</sup> +1)
N-b-196	NA	N-b-171		Me	MeOC(O)	H	H	1Me-5-1Hldz	C		370 (M <sup>+</sup> +1)
N-b-197	NF-NA	N-b-1	BzCl	Me	Bz	H	H	2-Nap	C		396 (M <sup>+</sup> +1)
N-b-198	NF-NA	N-b-3	BzCl	Me	Bz	H	H	5-Ind	C		399 (M <sup>+</sup> +1)
N-b-199	NF-NA	N-b-5	BzCl	Me	Bz	H	H	1Me-5-Ind	C		399 (M <sup>+</sup> +1)
N-b-200	NF-NA	N-b-9	BzCl	Me	Bz	H	H	5-1Hldz	C		386 (M <sup>+</sup> +1)
N-b-201	NF-NA	N-b-11	BzCl	Me	Bz	H	H	1Me-5-1Hldz	C		400 (M <sup>+</sup> +1)
N-b-202	NF-NA	N-b-1	PhOCOCi	Me		H	H	2-Nap	C		412 (M <sup>+</sup> +1)
N-b-203	NF-NA	N-b-5	PhOCOCi	Me		H	H	1Me-5-Ind	C		415 (M <sup>+</sup> +1)
N-b-204	NF-NA	N-b-1	cPenCH2COCi	Me		H	H	2-Nap	C		402 (M <sup>+</sup> +1)
N-b-205	NF-NA	N-b-3	cPenCH2COCi	Me		H	H	1Me-5-Ind	C		405 (M <sup>+</sup> +1)
N-b-206	NF-NA	N-b-1		Me		H	H	2-Nap	C		403 (M <sup>+</sup> +1)
N-b-207	NF-NA	N-b-5		Me		H	H	1Me-5-Ind	C		406 (M <sup>+</sup> +1)
N-b-208	NF-NA	N-b-1	PhNCO	Me	PhNHC(O)	H	H	2-Nap	C		411 (M <sup>+</sup> +1)
N-b-209	NF-NA	N-b-5	PhNCO	Me	PhNHC(O)	H	H	1Me-5-Ind	C		414 (M <sup>+</sup> +1)
N-b-210	NF-NA	N-b-1	cHexNCO	Me	cHexNHC(O)	H	H	2-Nap	C		417 (M <sup>+</sup> +1)
N-b-211	NF-NA	N-b-5	cHexNCO	Me	cHexNHC(O)	H	H	1Me-5-Ind	C		420 (M <sup>+</sup> +1)
N-b-212	NF-NA	N-b-1	cHexNCS	Me	PhNHC(S)	H	H	2-Nap	C		430 (M <sup>+</sup> +1)

[Example N-c-51]

Synthesis of ethyl 3-[4-(imidazol-1-yl)-3-(naphthalen-2-yl)phenyl]acrylate

(Compound No. N-c-51) (Synthesis method NB1)

According to the procedure described in the synthesis method of the

compound of Example N-a-1 (Synthesis method NB1) provided that the reaction was carried out for 16 hours, and the column chromatography was performed with chloroform:methanol =100:1, Intermediate n-33 (300.4 mg), 2-naphthaleneboronic acid (208.3 mg), 2 M aqueous sodium carbonate (900  $\mu$ l) and (Ph<sub>3</sub>P)<sub>4</sub>Pd (108.3 mg) were reacted and treated to obtain the title compound (Intermediate N-c-51, 304.2 mg).

[Example N-c-52]

Synthesis of 3-[4-(imidazol-1-yl)-3-(naphthalen-2-yl)phenyl]acrylic acid (Compound No. N-c-51) (Synthesis method NA)

According to the procedure described in the synthesis method of the compound of Example N-a-2 (Synthesis method NA) provided that the reaction was carried out for 2 hours, the compound of Example N-c-51 (301.2 mg) and 2 N aqueous sodium hydroxide (980  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. N-c-52, 286.4 mg).

[Examples N-c-1 to N-c-64]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification including the examples described above are shown in Table-N-C-1 to Table-N-C-3. In the tables, the compound numbers are mentioned in the columns indicated as "Exp.". In the tables, used methods among the aforementioned synthesis methods are shown in the columns of "Syn" with symbols, the starting compounds 1 are mentioned in the columns of "SM1", and the starting compounds 2 are mentioned in the columns of "SM2".

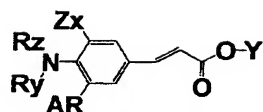


Table-N-C-1

Exp.	Syn	SM1	SM2	NRzRy	Y	Zx	AR	LCMS		
								method	RTime	Mass
N-c-1	NB1	Int.n-36	BRA1		Me	H	2-Nap	C		358 (M <sup>+</sup> +1)
N-c-2	NA	N-c-1			H	H	2-Nap	C		344 (M <sup>+</sup> +1)
N-c-3	NB1	Int.n-36	BRA2		Me	H	5-Ind	C		347 (M <sup>+</sup> +1)
N-c-4	NA	N-c-3			H	H	5-Ind	C		333 (M <sup>+</sup> +1)
N-c-5	NB1	Int.n-36	BRA3		Me	H	1Me-5-Ind	C		361 (M <sup>+</sup> +1)
N-c-6	NA	N-c-5			H	H	1Me-5-Ind	C		347 (M <sup>+</sup> +1)
N-c-7	NB1	Int.n-36	BRA5		Me	H	5-1HIdz	C		348 (M <sup>+</sup> +1)
N-c-8	NA	N-c-7			H	H	5-1HIdz	C		334 (M <sup>+</sup> +1)
N-c-9	NB1	Int.n-36	BRA6		Me	H	1Me-5-1HIdz	C		362 (M <sup>+</sup> +1)
N-c-10	NA	N-c-9			H	H	1Me-5-1HIdz	C		348 (M <sup>+</sup> +1)
N-c-11	NB1	Int.n-36	BRA9		Me	H	5-Bzt	C		365 (M <sup>+</sup> +1)
N-c-12	NA	N-c-11			H	H	5-Bzt	C		351 (M <sup>+</sup> +1)
N-c-13	NB1	Int.n-36	BRA10		Me	H	3-Qu	C		359 (M <sup>+</sup> +1)
N-c-14	NA	N-c-13			H	H	3-Qu	C		345 (M <sup>+</sup> +1)
N-c-15	NB1	Int.n-36	BRA11		Me	H	6-Qu	C		359 (M <sup>+</sup> +1)
N-c-16	NA	N-c-15			H	H	6-Qu	C		345 (M <sup>+</sup> +1)
N-c-17	NB1	Int.n-37	BRA1		Me	H	2-Nap	C		374 (M <sup>+</sup> +1)
N-c-18	NA	N-c-17			H	H	2-Nap	C		360 (M <sup>+</sup> +1)
N-c-19	NB1	Int.n-37	BRA2		Me	H	5-Ind	C		363 (M <sup>+</sup> +1)
N-c-20	NA	N-c-19			H	H	5-Ind	C		349 (M <sup>+</sup> +1)
N-c-21	NB1	Int.n-37	BRA3		Me	H	1Me-5-Ind	C		377 (M <sup>+</sup> +1)
N-c-22	NA	N-c-21			H	H	1Me-5-Ind	C		363 (M <sup>+</sup> +1)



Table-N-C-2











































Exp.	Syn.	SM1	SM2	NRzRy	Y	Zx	AR	LCMS		
								method	RTime	Mass
N-c-23	NB1	Int.n-37	BRA5		Me	H	5-1HIdz	C		364 (M <sup>+</sup> +1)
N-c-24	NA	N-c-23			H	H	5-1HIdz	C		350 (M <sup>+</sup> +1)
N-c-25	NB1	Int.n-37	BRA6		Me	H	1Me-5-1HIdz	C		378 (M <sup>+</sup> +1)
N-c-26	NA	N-c-25			H	H	1Me-5-1HIdz	C		364 (M <sup>+</sup> +1)
N-c-27	NB1	Int.n-26	BRA1		Me	H	2-Nap	C		372 (M <sup>+</sup> +1)
N-c-28	NA	N-c-27			H	H	2-Nap	C		358 (M <sup>+</sup> +1)
N-c-29	NB1	Int.n-26	BRA2		Me	H	5-Ind	C		361 (M <sup>+</sup> +1)
N-c-30	NA	N-c-29			H	H	5-Ind	C		347 (M <sup>+</sup> +1)
N-c-31	NB1	Int.n-26	BRA3		Me	H	1Me-5-Ind	C		375 (M <sup>+</sup> +1)
N-c-32	NA	N-c-31			H	H	1Me-5-Ind	C		361 (M <sup>+</sup> +1)
N-c-33	NB1	Int.n-26	BRA5		Me	H	5-1HIdz	C		362 (M <sup>+</sup> +1)
N-c-34	NA	N-c-33			H	H	5-1HIdz	C		348 (M <sup>+</sup> +1)
N-c-35	NB1	Int.n-26	BRA6		Me	H	1Me-5-1HIdz	C		376 (M <sup>+</sup> +1)
N-c-36	NA	N-c-35			H	H	1Me-5-1HIdz	C		362 (M <sup>+</sup> +1)
N-c-37	NB1	Int.n-28	BRA1		Me	H	2-Nap	C		386 (M <sup>+</sup> +1)
N-c-38	NA	N-c-37			H	H	2-Nap	C		372 (M <sup>+</sup> +1)
N-c-39	NB1	Int.n-28	BRA3		Me	H	1Me-5-Ind	C		389 (M <sup>+</sup> +1)
N-c-40	NA	N-c-39			H	H	1Me-5-Ind	C		375 (M <sup>+</sup> +1)
N-c-41	NB1	Int.n-28	BRA5		Me	H	5-1HIdz	C		376 (M <sup>+</sup> +1)
N-c-42	NA	N-c-41			H	H	5-1HIdz	C		362 (M <sup>+</sup> +1)
N-c-43	NB1	Int.n-28	BRA6		Me	H	1Me-5-1HIdz	C		390 (M <sup>+</sup> +1)
N-c-44	NA	N-c-43			H	H	1Me-5-1HIdz	C		376 (M <sup>+</sup> +1)

Table-N-C-3

Exp.	Syn	SM1	SM2	NRzRy	Y	Zx	AR	LCMS		
								method	RTime	Mass
N-c-45	NB1	Int.n-30	BRA3		Me	H	1Me-5-Ind	C		389 (M <sup>+</sup> +1)
N-c-46	NA	N-c-45			H	H	1Me-5-Ind	C		375 (M <sup>+</sup> +1)
N-c-47	NB1	Int.n-30	BRA5		Me	H	5-1HIdz	C		376 (M <sup>+</sup> +1)
N-c-48	NA	N-c-47			H	H	5-1HIdz	C		362 (M <sup>+</sup> +1)
N-c-49	NB1	Int.n-30	BRA6		Me	H	1Me-5-1HIdz	C		390 (M <sup>+</sup> +1)
N-c-50	NA	N-c-49			H	H	1Me-5-1HIdz	C		376 (M <sup>+</sup> +1)
N-c-51	NB1	Int.n-33	BRA1		Et	H	2-Nap	C		369 (M <sup>+</sup> +1)
N-c-52	NA	N-c-51			H	H	2-Nap	C		341 (M <sup>+</sup> +1)
N-c-53	NB1	Int.n-33	BRA3		Et	H	1Me-5-Ind	C		372 (M <sup>+</sup> +1)
N-c-54	NA	N-c-53			H	H	1Me-5-Ind	C		344 (M <sup>+</sup> +1)
N-c-55	NB1	Int.n-33	BRA6		Et	H	1Me-5-1HIdz	C		373 (M <sup>+</sup> +1)
N-c-56	NA	N-c-55			H	H	1Me-5-1HIdz	C		345 (M <sup>+</sup> +1)
N-c-57	NB1	Int.n-35	BRA1		Et	H	2-Nap	C		368 (M <sup>+</sup> +1)
N-c-58	NA	N-c-57			H	H	2-Nap	C		340 (M <sup>+</sup> +1)
N-c-59	NB1	Int.n-35	BRA3		Et	H	1Me-5-Ind	C		371 (M <sup>+</sup> +1)
N-c-60	NA	N-c-59			H	H	1Me-5-Ind	C		343 (M <sup>+</sup> +1)
N-c-61	NB1	Int.n-35	BRA5		Et	H	5-1HIdz	C		358 (M <sup>+</sup> +1)
N-c-62	NA	N-c-61			H	H	5-1HIdz	C		330 (M <sup>+</sup> +1)
N-c-63	NB1	Int.n-35	BRA6		Et	H	1Me-5-1HIdz	C		372 (M <sup>+</sup> +1)
N-c-64	NA	N-c-63			H	H	1Me-5-1HIdz	C		344 (M <sup>+</sup> +1)

[Example N-d-61]

Synthesis of ethyl 3-[4-(imidazol-1-yl)-3-(naphthalen-2-yl)phenyl]propionate

(Compound No. N-d-51) (Synthesis method ND1)

According to the procedure described in the synthesis method of

Intermediate n-7 (Synthesis method ND1) provided that the reaction was carried

out for 6 hours, the compound of Example N-c-51 (301.5 mg) and 10% palladium/carbon (67.3 mg) were reacted and treated to obtain the title compound (Compound No. N-d-61, 143.5 mg).

[Example N-d-62]

Synthesis of 3-[4-(imidazol-1-yl)-3-(naphthalen-2-yl)phenyl]propionic acid (Compound No. N-d-62) (Synthesis method NA)

According to the procedure described in the synthesis method of the compound of Example N-a-2 (Synthesis method NA) provided that the reaction was carried out for 3 hours, the compound of Example N-d-61 (140.3 mg) and 2 N aqueous sodium hydroxide (600  $\mu$ l) were reacted and treated to obtain the title compound (Compound No. N-d-62, 100.4 mg).

[Examples N-d-1 to N-d-74]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification including the examples described above are shown in Table-N-D-1 to Table-N-D-4. In the tables, the compound numbers are mentioned in the columns indicated as "Exp.". In the tables, used methods among the aforementioned synthesis methods are shown in the columns of "Syn" with symbols, the starting compounds 1 are mentioned in the columns of "SM1", and the starting compounds 2 are mentioned in the columns of "SM2".

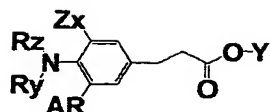


Table-N-D-1

Exp.	Syn	SM1	SM2	NRzRy	Y	Zx	AR	- LCMS		
								method	RTime	Mass
N-d-1	NB1	Int.n-21	BRA1		Me	H	2-Nap	C		360 (M <sup>+</sup> +1)
N-d-2	NA	N-d-1			H	H	2-Nap	C		346 (M <sup>+</sup> +1)
N-d-3	NB1	Int.n-21	BRA2		Me	H	5-Ind	D	4.79	349 (M <sup>+</sup> +1)
N-d-4	NA	N-d-3			H	H	5-Ind	D	3.54	335 (M <sup>+</sup> +1)
N-d-5	NB1	Int.n-21	BRA3		Me	H	1Me-5-Ind	D	5.72	363 (M <sup>+</sup> +1)
N-d-6	NA	N-d-5			H	H	1Me-5-Ind	D	4.31	349 (M <sup>+</sup> +1)
N-d-7	NB1	Int.n-21	BRA5		Me	H	5-1HIdz	C		350 (M <sup>+</sup> +1)
N-d-8	NA	N-d-7			H	H	5-1HIdz	C		336 (M <sup>+</sup> +1)
N-d-9	NB1	Int.n-21	BRA6		Me	H	1Me-5-1HIdz	C		364 (M <sup>+</sup> +1)
N-d-10	NA	N-d-9			H	H	1Me-5-1HIdz	C		350 (M <sup>+</sup> +1)
N-d-11	NB1	Int.n-21	BRA9		Me	H	5-Bzt	C		367 (M <sup>+</sup> +1)
N-d-12	NA	N-d-11			H	H	5-Bzt	C		353 (M <sup>+</sup> +1)
N-d-13	NB1	Int.n-21	BRA10		Me	H	3-Qu	C		361 (M <sup>+</sup> +1)
N-d-14	NA	N-d-13			H	H	3-Qu	C		347 (M <sup>+</sup> +1)
N-d-15	NB1	Int.n-21	BRA11		Me	H	6-Qu	C		361 (M <sup>+</sup> +1)
N-d-16	NA	N-d-15			H	H	6-Qu	C		347 (M <sup>+</sup> +1)
N-d-17	NB1	Int.n-24	BRA1		Me	H	2-Nap	C		376 (M <sup>+</sup> +1)
N-d-18	NA	N-d-17			H	H	2-Nap	C		362 (M <sup>+</sup> +1)
N-d-19	NB1	Int.n-24	BRA2		Me	H	5-Ind	C		365 (M <sup>+</sup> +1)
N-d-20	NA	N-d-19			H	H	5-Ind	C		351 (M <sup>+</sup> +1)
N-d-21	NB1	Int.n-24	BRA3		Me	H	1Me-5-Ind	C		379 (M <sup>+</sup> +1)
N-d-22	NA	N-d-21			H	H	1Me-5-Ind	C		365 (M <sup>+</sup> +1)

Table-N-D-2























Exp.	Syn	SM1	SM2	NRzRy	Y	Zx	AR	LCMS		
								method	RTime	Mass
N-d-23	NB1	Int.n-24	BRA5		Me	H	5-1HIdz	C		366 (M <sup>+</sup> +1)
N-d-24	NA	N-d-23			H	H	5-1HIdz	C		352 (M <sup>+</sup> +1)
N-d-25	NB1	Int.n-24	BRA6		Me	H	1Me-5-1HIdz	C		380 (M <sup>+</sup> +1)
N-d-26	NA	N-d-25			H	H	1Me-5-1HIdz	C		366 (M <sup>+</sup> +1)
N-d-27	NB1	Int.n-24	BRA9		Me	H	5-Bzt	C		383 (M <sup>+</sup> +1)
N-d-28	NA	N-d-27			H	H	5-Bzt	C		369 (M <sup>+</sup> +1)
N-d-29	NB1	Int.n-24	BRA11		Me	H	6-Qu	C		377 (M <sup>+</sup> +1)
N-d-30	NA	N-d-29			H	H	6-Qu	C		363 (M <sup>+</sup> +1)
N-d-31	NB1	Int.n-27	BRA1		Me	H	2-Nap	C		374 (M <sup>+</sup> +1)
N-d-32	NA	N-d-31			H	H	2-Nap	C		360 (M <sup>+</sup> +1)
N-d-33	NB1	Int.n-27	BRA2		Me	H	5-Ind	C		363 (M <sup>+</sup> +1)
N-d-34	NA	N-d-33			H	H	5-Ind	C		349 (M <sup>+</sup> +1)
N-d-35	NB1	Int.n-27	BRA3		Me	H	1Me-5-Ind	C		377 (M <sup>+</sup> +1)
N-d-36	NA	N-d-35			H	H	1Me-5-Ind	C		363 (M <sup>+</sup> +1)
N-d-37	NB1	Int.n-27	BRA5		Me	H	5-1HIdz	C		364 (M <sup>+</sup> +1)
N-d-38	NA	N-d-37			H	H	5-1HIdz	C		350 (M <sup>+</sup> +1)
N-d-39	NB1	Int.n-27	BRA6		Me	H	1Me-5-1HIdz	C		378 (M <sup>+</sup> +1)
N-d-40	NA	N-d-39			H	H	1Me-5-1HIdz	C		364 (M <sup>+</sup> +1)
N-d-41	NB1	Int.n-27	BRA11		Me	H	6-Qu	C		375 (M <sup>+</sup> +1)
N-d-42	NA	N-d-41			H	H	6-Qu	C		361 (M <sup>+</sup> +1)
N-d-43	NB1	Int.n-27	BRA9		Me	H	5-Bzt	C		381 (M <sup>+</sup> +1)
N-d-44	NA	N-d-43			H	H	5-Bzt	C		367 (M <sup>+</sup> +1)

Table-N-D-3

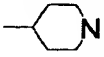
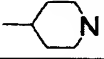

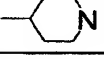
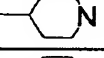

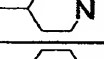
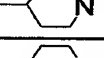
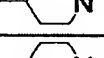
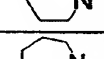




















Exp.	Syn	SM1	SM2	NRzRy	Y	Zx	AR	LCMS		
								method	RTime	Mass
N-d-45	NB1	Int.n-29	BRA1		Me	H	2-Nap	C		388 (M <sup>+</sup> +1)
N-d-46	NA	N-d-45			H	H	2-Nap	C		374 (M <sup>+</sup> +1)
N-d-47	NB1	Int.n-29	BRA3		Me	H	1Me-5-Ind	C		391 (M <sup>+</sup> +1)
N-d-48	NA	N-d-47			H	H	1Me-5-Ind	C		377 (M <sup>+</sup> +1)
N-d-49	NB1	Int.n-29	BRA5		Me	H	5-1Idz	C		378 (M <sup>+</sup> +1)
N-d-50	NA	N-d-49			H	H	5-1Idz	C		364 (M <sup>+</sup> +1)
N-d-51	NB1	Int.n-29	BRA6		Me	H	1Me-5-1HIdz	C		392 (M <sup>+</sup> +1)
N-d-52	NA	N-d-51			H	H	1Me-5-1HIdz	C		378 (M <sup>+</sup> +1)
N-d-53	NB1	Int.n-29	BRA10		Me	H	3-Qu	C		389 (M <sup>+</sup> +1)
N-d-54	NA	N-d-53			H	H	3-Qu	C		375 (M <sup>+</sup> +1)
N-d-55	NB1	Int.n-31	BRA3		Me	H	1Me-5-Ind	C		391 (M <sup>+</sup> +1)
N-d-56	NA	N-d-55			H	H	1Me-5-Ind	C		377 (M <sup>+</sup> +1)
N-d-57	NB1	Int.n-31	BRA5		Me	H	5-1Idz	C		378 (M <sup>+</sup> +1)
N-d-58	NA	N-d-57			H	H	5-1Idz	C		364 (M <sup>+</sup> +1)
N-d-59	NB1	Int.n-31	BRA6		Me	H	1Me-5-1HIdz	C		392 (M <sup>+</sup> +1)
N-d-60	NA	N-d-59			H	H	1Me-5-1HIdz	C		378 (M <sup>+</sup> +1)
N-d-61	ND1	N-c-51			Et	H	2-Nap	C		371 (M <sup>+</sup> +1)
N-d-62	NA	N-d-61			H	H	2-Nap	C		343 (M <sup>+</sup> +1)
N-d-63	ND1	N-c-53			Et	H	1Me-5-Ind	C		374 (M <sup>+</sup> +1)
N-d-64	NA	N-d-63			H	H	1Me-5-Ind	C		346 (M <sup>+</sup> +1)
N-d-65	ND1	N-c-55			Et	H	1Me-5-1HIdz	C		375 (M <sup>+</sup> +1)
N-d-66	NA	N-d-65			H	H	1Me-5-1HIdz	C		347 (M <sup>+</sup> +1)

Table-N-D-4

Exp.	Syn.	SM1	SM2	NRzRy	Y	Zx	AR	LCMS		
								method	RTime	Mass
N-d-67	ND1	N-c-57			Et	H	2-Nap	C		370 (M <sup>+</sup> +1)
N-d-68	NA	N-d-45			H	H	2-Nap	C		342 (M <sup>+</sup> +1)
N-d-69	ND1	N-c-59			Et	H	1Me-5-Ind	C		373 (M <sup>+</sup> +1)
N-d-70	NA	N-d-47			H	H	1Me-5-Ind	C		345 (M <sup>+</sup> +1)
N-d-71	ND1	N-c-61			Et	H	5-1Idz	C		360 (M <sup>+</sup> +1)
N-d-72	NA	N-d-49			H	H	5-1Idz	C		332 (M <sup>+</sup> +1)
N-d-73	ND1	N-c-63			Et	H	1Me-5-1HIdz	C		374 (M <sup>+</sup> +1)
N-d-74	NA	N-d-51			H	H	1Me-5-1HIdz	C		346 (M <sup>+</sup> +1)

[Examples N-e-1 to N-e-204]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification are shown in Table-N-E-1 to Table-N-E-7. In the tables, the compound numbers are mentioned in the columns indicated as "Exp.". In the tables, corresponding methods among the aforementioned synthesis methods are shown in the columns of "Syn" with symbols, the starting compounds 1 are mentioned in the columns of "SM1", and the starting compounds 2 are mentioned in the columns of "SM2".

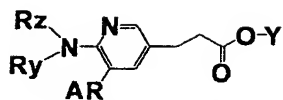


Table-N-E-1

Exp.	Syn	SM1	SM2	NRzRy	Y	AR	LCMS		
							method	RTime	Mass
N-e-1	NB1	Int.n-48	BRA1		Et	2-Nap	C		389 (M <sup>+</sup> +1)
N-e-2	NA	N-e-1			H	2-Nap	C		375 (M <sup>+</sup> +1)
N-e-3	NB1	Int.n-48	BRA2		Et	5-Ind	C		378 (M <sup>+</sup> +1)
N-e-4	NA	N-e-3			H	5-Ind	C		364 (M <sup>+</sup> +1)
N-e-5	NB1	Int.n-48	BRA3		Et	1Me-5-Ind	C		392 (M <sup>+</sup> +1)
N-e-6	NA	N-e-5			H	1Me-5-Ind	C		378 (M <sup>+</sup> +1)
N-e-7	NB1	Int.n-48	BRA5		Et	5-1HIdz	C		379 (M <sup>+</sup> +1)
N-e-8	NA	N-e-7			H	5-1HIdz	C		365 (M <sup>+</sup> +1)
N-e-9	NB1	Int.n-48	BRA6		Et	1Me-5-1HIdz	C		393 (M <sup>+</sup> +1)
N-e-10	NA	N-e-9			H	1Me-5-1HIdz	C		379 (M <sup>+</sup> +1)
N-e-11	NB1	Int.n-48	BRA10		Et	3-Qu	C		390 (M <sup>+</sup> +1)
N-e-12	NA	N-e-11			H	3-Qu	C		376 (M <sup>+</sup> +1)
N-e-13	NB1	Int.n-48	BRA11		Et	6-Qu	C		390 (M <sup>+</sup> +1)
N-e-14	NA	N-e-13			H	6-Qu	C		376 (M <sup>+</sup> +1)
N-e-15	NB1	Int.n-48	BRA12		Et	6-IQ	C		390 (M <sup>+</sup> +1)
N-e-16	NA	N-e-15			H	6-IQ	C		376 (M <sup>+</sup> +1)
N-e-17	NB1	Int.n-49	BRA1		Et	2Nap	C		375 (M <sup>+</sup> +1)
N-e-18	NA	N-e-17			H	2Nap	C		361 (M <sup>+</sup> +1)
N-e-19	NB1	Int.n-49	BRA2		Et	5-Ind	C		364 (M <sup>+</sup> +1)
N-e-20	NA	N-e-19			H	5-Ind	C		350 (M <sup>+</sup> +1)
N-e-21	NB1	Int.n-49	BRA3		Et	1Me-5-Ind	C		378 (M <sup>+</sup> +1)
N-e-22	NA	N-e-21			H	1Me-5-Ind	C		364 (M <sup>+</sup> +1)



Table-N-E-2































Exp.	Syn	SM1	SM2	NRzRy	Y	AR	LCMS		
							method	RTime	Mass
N-e-23	NB1	Int.n-49	BRA5		Et	5-1HIdz	C		365 (M <sup>+</sup> +1)
N-e-24	NA	N-e-23			H	5-1HIdz	C		351 (M <sup>+</sup> +1)
N-e-25	NB1	Int.n-49	BRA6		Et	1Me-5-1HIdz	C		379 (M <sup>+</sup> +1)
N-e-26	NA	N-e-25			H	1Me-5-1HIdz	C		365 (M <sup>+</sup> +1)
N-e-27	NB1	Int.n-50	BRA1		Et	2-Nap	C		403 (M <sup>+</sup> +1)
N-e-28	NA	N-e-27			H	2-Nap	C		389 (M <sup>+</sup> +1)
N-e-29	NB1	Int.n-50	BRA2		Et	5-Ind	C		392 (M <sup>+</sup> +1)
N-e-30	NA	N-e-29			H	5-Ind	C		378 (M <sup>+</sup> +1)
N-e-31	NB1	Int.n-50	BRA3		Et	1Me-5-Ind	C		406 (M <sup>+</sup> +1)
N-e-32	NA	N-e-31			H	1Me-5-Ind	C		392 (M <sup>+</sup> +1)
N-e-33	NB1	Int.n-50	BRA5		Et	5-1HIdz	C		393 (M <sup>+</sup> +1)
N-e-34	NA	N-e-33			H	5-1HIdz	C		379 (M <sup>+</sup> +1)
N-e-35	NB1	Int.n-50	BRA6		Et	1Me-5-1HIdz	C		407 (M <sup>+</sup> +1)
N-e-36	NA	N-e-35			H	1Me-5-1HIdz	C		393 (M <sup>+</sup> +1)
N-e-37	NB1	Int.n-51	BRA1		Et	2-Nap	C		391 (M <sup>+</sup> +1)
N-e-38	NA	N-e-37			H	2-Nap	C		377 (M <sup>+</sup> +1)
N-e-39	NB1	Int.n-51	BRA3		Et	1Me-5-Ind	C		394 (M <sup>+</sup> +1)
N-e-40	NA	N-e-39			H	1Me-5-Ind	C		380 (M <sup>+</sup> +1)
N-e-41	NB1	Int.n-51	BRA5		Et	5-1HIdz	C		381 (M <sup>+</sup> +1)
N-e-42	NA	N-e-41			H	5-1HIdz	C		367 (M <sup>+</sup> +1)
N-e-43	NB1	Int.n-51	BRA6		Et	1Me-5-1HIdz	C		395 (M <sup>+</sup> +1)
N-e-44	NA	N-e-43			H	1Me-5-1HIdz	C		381 (M <sup>+</sup> +1)

Table-N-E-3

Exp.	Syn	SM1	SM2	NRzRy	Y	AR	LCMS		
							method	RTime	Mass
N-e-45	NB1	Intn-52	BRA1	 N	Et	2-Nap	C		403 (M <sup>+</sup> +1)
N-e-46	NA	N-e-45		 N	H	2-Nap	C		389 (M <sup>+</sup> +1)
N-e-47	NB1	Intn-52	BRA3	 N	Et	1Me-5-Ind	C		406 (M <sup>+</sup> +1)
N-e-48	NA	N-e-47		 N	H	1Me-5-Ind	C		392 (M <sup>+</sup> +1)
N-e-49	NB1	Intn-52	BRA5	 N	Et	5-1HIdz	C		393 (M <sup>+</sup> +1)
N-e-50	NA	N-e-49		 N	H	5-1HIdz	C		379 (M <sup>+</sup> +1)
N-e-51	NB1	Intn-52	BRA6	 N	Et	1Me-5-1HIdz	C		407 (M <sup>+</sup> +1)
N-e-52	NA	N-e-51		 N	H	1Me-5-1HIdz	C		393 (M <sup>+</sup> +1)

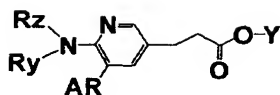


Table-N-E-4

Exp.	Syn	SM1	SM2	Rz	Ry	Y	AR	LCMS		
								method	RTime	Mass
N-e-53	NB1	Int.n-53	BRA1	Et	Me	Et	2-Nap	C		363 (M <sup>+</sup> +1)
N-e-54	NA	N-e-53		Et	Me	H	2-Nap	C		335 (M <sup>+</sup> +1)
N-e-55	NB1	Int.n-53	BRA2	Et	Me	Et	5-Ind	C		352 (M <sup>+</sup> +1)
N-e-56	NA	N-e-55		Et	Me	H	5-Ind	C		324 (M <sup>+</sup> +1)
N-e-57	NB1	Int.n-53	BRA3	Et	Me	Et	1Me-5-Ind	C		366 (M <sup>+</sup> +1)
N-e-58	NA	N-e-57		Et	Me	H	1Me-5-Ind	C		338 (M <sup>+</sup> +1)
N-e-59	NB1	Int.n-53	BRA5	Et	Me	Et	5-1HIdz	C		353 (M <sup>+</sup> +1)
N-e-60	NA	N-e-59		Et	Me	H	5-1HIdz	C		325 (M <sup>+</sup> +1)
N-e-61	NB1	Int.n-53	BRA6	Et	Me	Et	1Me-5-1HIdz	C		367 (M <sup>+</sup> +1)
N-e-62	NA	N-e-61		Et	Me	H	1Me-5-1HIdz	C		339 (M <sup>+</sup> +1)
N-e-63	NB1	Int.n-54	BRA1	Et	Et	Et	2-Nap	C		377 (M <sup>+</sup> +1)
N-e-64	NA	N-b-63		Et	Et	H	2-Nap	C		349 (M <sup>+</sup> +1)
N-e-65	NB1	Int.n-54	BRA2	Et	Et	Et	5-Ind	C		366 (M <sup>+</sup> +1)
N-e-66	NA	N-b-65		Et	Et	H	5-Ind	C		338 (M <sup>+</sup> +1)
N-e-67	NB1	Int.n-54	BRA3	Et	Et	Et	1Me-5-Ind	C		380 (M <sup>+</sup> +1)
N-e-68	NA	N-b-67		Et	Et	H	1Me-5-Ind	C		352 (M <sup>+</sup> +1)
N-e-69	NB1	Int.n-54	BRA5	Et	Et	Et	5-1HIdz	C		367 (M <sup>+</sup> +1)
N-e-70	NA	N-b-69		Et	Et	H	5-1HIdz	C		339 (M <sup>+</sup> +1)
N-e-71	NB1	Int.n-54	BRA6	Et	Et	Et	1Me-5-1HIdz	C		381 (M <sup>+</sup> +1)
N-e-72	NA	N-b-71		Et	Et	H	1Me-5-1HIdz	C		353 (M <sup>+</sup> +1)
N-e-73	NB1	Int.n-55	BRA1	nPr	Me	Et	2-Nap	C		377 (M <sup>+</sup> +1)
N-e-74	NA	N-b-73		nPr	Me	H	2-Nap	C		349 (M <sup>+</sup> +1)
N-e-75	NB1	Int.n-55	BRA2	nPr	Me	Et	5-Ind	C		366 (M <sup>+</sup> +1)
N-e-76	NA	N-b-75		nPr	Me	H	5-Ind	C		338 (M <sup>+</sup> +1)
N-e-77	NB1	Int.n-55	BRA3	nPr	Me	Et	1Me-5-Ind	C		380 (M <sup>+</sup> +1)
N-e-78	NA	N-b-77		nPr	Me	H	1Me-5-Ind	C		352 (M <sup>+</sup> +1)
N-e-79	NB1	Int.n-55	BRA5	nPr	Me	Et	5-1HIdz	C		367 (M <sup>+</sup> +1)
N-e-80	NA	N-b-79		nPr	Me	H	5-1HIdz	C		339 (M <sup>+</sup> +1)
N-e-81	NB1	Int.n-55	BRA6	nPr	Me	Et	1Me-5-1HIdz	C		381 (M <sup>+</sup> +1)
N-e-82	NA	N-b-81		nPr	Me	H	1Me-5-1HIdz	C		353 (M <sup>+</sup> +1)
N-e-83	NB1	Int.n-56	BRA1	iPr	Me	Et	2-Nap	C		377 (M <sup>+</sup> +1)
N-e-84	NA	N-b-83		iPr	Me	H	2-Nap	C		349 (M <sup>+</sup> +1)
N-e-85	NB1	Int.n-56	BRA2	iPr	Me	Et	5-Ind	C		366 (M <sup>+</sup> +1)
N-e-86	NA	N-b-85		iPr	Me	H	5-Ind	C		338 (M <sup>+</sup> +1)
N-e-87	NB1	Int.n-56	BRA3	iPr	Me	Et	1Me-5-Ind	C		380 (M <sup>+</sup> +1)
N-e-88	NA	N-b-87		iPr	Me	H	1Me-5-Ind	C		352 (M <sup>+</sup> +1)
N-e-89	NB1	Int.n-56	BRA5	iPr	Me	Et	5-1HIdz	C		367 (M <sup>+</sup> +1)
N-e-90	NA	N-b-89		iPr	Me	H	5-1HIdz	C		339 (M <sup>+</sup> +1)
N-e-91	NB1	Int.n-56	BRA6	iPr	Me	Et	1Me-5-1HIdz	C		381 (M <sup>+</sup> +1)
N-e-92	NA	N-b-91		iPr	Me	H	1Me-5-1HIdz	C		353 (M <sup>+</sup> +1)
N-e-93	NB1	Int.n-57	BRA1	nBu	Me	Et	2-Nap	C		391 (M <sup>+</sup> +1)
N-e-94	NA	N-b-93		nBu	Me	H	2-Nap	C		363 (M <sup>+</sup> +1)
N-e-95	NB1	Int.n-57	BRA2	nBu	Me	Et	5-Ind	C		380 (M <sup>+</sup> +1)
N-e-96	NA	N-b-95		nBu	Me	H	5-Ind	C		352 (M <sup>+</sup> +1)

Table-N-E-5

Exp.	Syn	SM1	SM2	Rz	Ry	Y	AR	LCMS		
								method	RTime	Mass
N-e-97	NB1	Int.n-57	BRA3	nBu	Me	Et	1Me-5-Ind	C		394 (M <sup>+</sup> +1)
N-e-98	NA	N-e-97		nBu	Me	H	1Me-5-Ind	C		366 (M <sup>+</sup> +1)
N-e-99	NB1	Int.n-57	BRA5	nBu	Me	Et	5-1HIdz	C		381 (M <sup>+</sup> +1)
N-e-100	NA	N-e-99		nBu	Me	H	5-1HIdz	C		353 (M <sup>+</sup> +1)
N-e-101	NB1	Int.n-57	BRA6	nBu	Me	Et	1Me-5-1HIdz	C		395 (M <sup>+</sup> +1)
N-e-102	NA	N-e-101		nBu	Me	H	1Me-5-1HIdz	C		367 (M <sup>+</sup> +1)
N-e-103	NB1	Int.n-58	BRA1	iBu	Me	Et	2-Nap	C		391 (M <sup>+</sup> +1)
N-e-104	NA	N-e-103		iBu	Me	H	2-Nap	C		363 (M <sup>+</sup> +1)
N-e-105	NB1	Int.n-58	BRA2	iBu	Me	Et	5-Ind	C		380 (M <sup>+</sup> +1)
N-e-106	NA	N-e-105		iBu	Me	H	5-Ind	C		352 (M <sup>+</sup> +1)
N-e-107	NB1	Int.n-58	BRA3	iBu	Me	Et	1Me-5-Ind	C		394 (M <sup>+</sup> +1)
N-e-108	NA	N-e-107		iBu	Me	H	1Me-5-Ind	C		366 (M <sup>+</sup> +1)
N-e-109	NB1	Int.n-58	BRA5	iBu	Me	Et	5-1HIdz	C		381 (M <sup>+</sup> +1)
N-e-110	NA	N-e-109		iBu	Me	H	5-1HIdz	C		353 (M <sup>+</sup> +1)
N-e-111	NB1	Int.n-58	BRA6	iBu	Me	Et	1Me-5-1HIdz	C		395 (M <sup>+</sup> +1)
N-e-112	NA	N-e-111		iBu	Me	H	1Me-5-1HIdz	C		367 (M <sup>+</sup> +1)
N-e-113	NB1	Int.n-62	BRA1	Bn	H	Et	2-Nap	C		411 (M <sup>+</sup> +1)
N-e-114	NA	N-e-113		Bn	H	H	2-Nap	C		383 (M <sup>+</sup> +1)
N-e-115	NB1	Int.n-62	BRA2	Bn	H	Et	5-Ind	C		400 (M <sup>+</sup> +1)
N-e-116	NA	N-e-115		Bn	H	H	5-Ind	C		372 (M <sup>+</sup> +1)
N-e-117	NB1	Int.n-62	BRA3	Bn	H	Et	1Me-5-Ind	C		414 (M <sup>+</sup> +1)
N-e-118	NA	N-e-117		Bn	H	H	1Me-5-Ind	C		386 (M <sup>+</sup> +1)
N-e-119	NB1	Int.n-62	BRA5	Bn	H	Et	5-1HIdz	C		401 (M <sup>+</sup> +1)
N-e-120	NA	N-e-119		Bn	H	H	5-1HIdz	C		373 (M <sup>+</sup> +1)
N-e-121	NB1	Int.n-62	BRA6	Bn	H	Et	1Me-5-1HIdz	C		415 (M <sup>+</sup> +1)
N-e-122	NA	N-e-121		Bn	H	H	1Me-5-1HIdz	C		387 (M <sup>+</sup> +1)
N-e-123	NB1	Int.n-63	BRA1	4MeBn	H	Et	2-Nap	C		425 (M <sup>+</sup> +1)
N-e-124	NA	N-e-123		4MeBn	H	H	2-Nap	C		397 (M <sup>+</sup> +1)
N-e-125	NB1	Int.n-63	BRA2	4MeBn	Me	Et	5-Ind	C		414 (M <sup>+</sup> +1)
N-e-126	NA	N-e-125		4MeBn	Me	H	5-Ind	C		386 (M <sup>+</sup> +1)
N-e-127	NB1	Int.n-63	BRA5	4MeBn	Me	Et	5-1HIdz	C		415 (M <sup>+</sup> +1)
N-e-128	NA	N-e-127		4MeBn	Me	H	5-1HIdz	C		387 (M <sup>+</sup> +1)
N-e-129	NB1	Int.n-64	BRA2	3MeBn	Me	Et	5-Ind	C		414 (M <sup>+</sup> +1)
N-e-130	NA	N-e-129		3MeBn	Me	H	5-Ind	C		386 (M <sup>+</sup> +1)
N-e-131	NB1	Int.n-64	BRA3	3MeBn	Me	Et	1Me-5-Ind	C		428 (M <sup>+</sup> +1)
N-e-132	NA	N-e-131		3MeBn	Me	H	1Me-5-Ind	C		400 (M <sup>+</sup> +1)
N-e-133	NB1	Int.n-64	BRA5	3MeBn	Me	Et	5-1HIdz	C		415 (M <sup>+</sup> +1)
N-e-134	NA	N-e-133		3MeBn	Me	H	5-1HIdz	C		387 (M <sup>+</sup> +1)
N-e-135	NB1	Int.n-65	BRA1	2MeBn	Me	Et	2-Nap	C		425 (M <sup>+</sup> +1)
N-e-136	NA	N-e-135		2MeBn	Me	H	2-Nap	C		397 (M <sup>+</sup> +1)
N-e-137	NB1	Int.n-65	BRA3	2MeBn	Me	Et	1Me-5-Ind	C		428 (M <sup>+</sup> +1)
N-e-138	NA	N-e-137		2MeBn	Me	H	1Me-5-Ind	C		400 (M <sup>+</sup> +1)
N-e-139	NB1	Int.n-65	BRA6	2MeBn	Me	Et	1Me-5-1HIdz	C		429 (M <sup>+</sup> +1)
N-e-140	NA	N-e-139		2MeBn	Me	H	1Me-5-1HIdz	C		401 (M <sup>+</sup> +1)

Taele-N-E-6

Exp.	Syn	SM1	SM2	Rz	Ry	Y	AR	LCMS		
								method	RTime	Mass
N-e-141	NB1	Int.n-66	BRA1	4FBn	H	Et	2-Nap	C		429 (M <sup>+</sup> +1)
N-e-142	NA	N-e-141		4FBn	H	H	2-Nap	C		401 (M <sup>+</sup> +1)
N-e-143	NB1	Int.n-66	BRA3	4FBn	H	Et	1Me-5-Ind	C		432 (M <sup>+</sup> +1)
N-e-144	NA	N-e-143		4FBn	H	H	1Me-5-Ind	C		404 (M <sup>+</sup> +1)
N-e-145	NB1	Int.n-66	BRA6	4FBn	H	Et	1Me-5-1HIdz	C		433 (M <sup>+</sup> +1)
N-e-146	NA	N-e-145		4FBn	H	H	1Me-5-1HIdz	C		405 (M <sup>+</sup> +1)
N-e-147	NB1	Int.n-67	BRA1	3FBn	H	Et	2-Nap	C		429 (M <sup>+</sup> +1)
N-e-148	NA	N-e-147		3FBn	H	H	2-Nap	C		401 (M <sup>+</sup> +1)
N-e-149	NB1	Int.n-67	BRA2	3FBn	H	Et	5-Ind	C		418 (M <sup>+</sup> +1)
N-e-150	NA	N-e-149		3FBn	H	H	5-Ind	C		390 (M <sup>+</sup> +1)
N-e-151	NB1	Int.n-67	BRA3	3FBn	H	Et	1Me-5-Ind	C		432 (M <sup>+</sup> +1)
N-e-152	NA	N-e-151		3FBn	H	H	1Me-5-Ind	C		404 (M <sup>+</sup> +1)
N-e-153	NB1	Int.n-68	BRA3	2FBn	H	Et	1Me-5-Ind	C		432 (M <sup>+</sup> +1)
N-e-154	NA	N-e-153		2FBn	H	H	1Me-5-Ind	C		404 (M <sup>+</sup> +1)
N-e-155	NB1	Int.n-68	BRA5	2FBn	H	Et	5-1HIdz	C		419 (M <sup>+</sup> +1)
N-e-156	NA	N-e-155		2FBn	H	H	5-1HIdz	C		391 (M <sup>+</sup> +1)
N-e-157	NB1	Int.n-68	BRA6	2FBn	H	Et	1Me-5-1HIdz	C		433 (M <sup>+</sup> +1)
N-e-158	NA	N-e-157		2FBn	H	H	1Me-5-1HIdz	C		405 (M <sup>+</sup> +1)
N-e-159	NB1	Int.n-69	BRA1	4MeOPh	H	Et	2-Nap	C		427 (M <sup>+</sup> +1)
N-e-160	NA	N-e-159		4MeOPh	H	H	2-Nap	C		399 (M <sup>+</sup> +1)
N-e-161	NB1	Int.n-69	BRA2	4MeOPh	H	Et	5-Ind	C		416 (M <sup>+</sup> +1)
N-e-162	NA	N-e-161		4MeOPh	H	H	5-Ind	C		388 (M <sup>+</sup> +1)
N-e-163	NB1	Int.n-69	BRA3	4MeOPh	H	Et	1Me-5-Ind	C		430 (M <sup>+</sup> +1)
N-e-164	NA	N-e-163		4MeOPh	H	H	1Me-5-Ind	C		402 (M <sup>+</sup> +1)
N-e-165	NB1	Int.n-69	BRA5	4MeOPh	H	Et	5-1HIdz	C		417 (M <sup>+</sup> +1)
N-e-166	NA	N-e-165		4MeOPh	H	H	5-1HIdz	C		389 (M <sup>+</sup> +1)
N-e-167	NB1	Int.n-70	BRA1	3MeOPh	H	Et	2-Nap	C		427 (M <sup>+</sup> +1)
N-e-168	NA	N-e-167		3MeOPh	H	H	2-Nap	C		399 (M <sup>+</sup> +1)
N-e-169	NB1	Int.n-70	BRA3	3MeOPh	H	Et	1Me-5-Ind	C		430 (M <sup>+</sup> +1)
N-e-170	NA	N-e-169		3MeOPh	H	H	1Me-5-Ind	C		402 (M <sup>+</sup> +1)
N-e-171	NB1	Int.n-70	BRA6	3MeOPh	H	Et	1Me-5-1HIdz	C		431 (M <sup>+</sup> +1)
N-e-172	NA	N-e-171		3MeOPh	H	H	1Me-5-1HIdz	C		403 (M <sup>+</sup> +1)
N-e-173	NB1	Int.n-71	BRA5	2MeOPh	H	Et	5-1HIdz	C		417 (M <sup>+</sup> +1)
N-e-174	NA	N-e-173		2MeOPh	H	H	5-1HIdz	C		389 (M <sup>+</sup> +1)
N-e-175	NB1	Int.n-71	BRA6	2MeOPh	H	Et	1Me-5-1HIdz	C		431 (M <sup>+</sup> +1)
N-e-176	NA	N-e-175		2MeOPh	H	H	1Me-5-1HIdz	C		403 (M <sup>+</sup> +1)
N-e-177	NB1	Int.n-71	BRA11	2MeOPh	H	Et	6-Qu	C		428 (M <sup>+</sup> +1)
N-e-178	NA	N-e-177		2MeOPh	H	H	6-Qu	C		400 (M <sup>+</sup> +1)
N-e-179	NB1	Int.n-72	BRA1	4CF3Ph	H	Et	2-Nap	C		465 (M <sup>+</sup> +1)
N-e-180	NA	N-e-179		4CF3Ph	H	H	2-Nap	C		437 (M <sup>+</sup> +1)
N-e-181	NB1	Int.n-72	BRA3	4CF3Ph	H	Et	1Me-5-Ind	C		468 (M <sup>+</sup> +1)
N-e-182	NA	N-e-181		4CF3Ph	H	H	1Me-5-Ind	C		440 (M <sup>+</sup> +1)
N-e-183	NB1	Int.n-72	BRA5	4CF3Ph	H	Et	5-1HIdz	C		455 (M <sup>+</sup> +1)
N-e-184	NA	N-e-183		4CF3Ph	H	H	5-1HIdz	C		427 (M <sup>+</sup> +1)
N-e-185	NB1	Int.n-72	BRA6	4CF3Ph	H	Et	1Me-5-1HIdz	C		469 (M <sup>+</sup> +1)
N-e-186	NA	N-e-185		4CF3Ph	H	H	1Me-5-1HIdz	C		441 (M <sup>+</sup> +1)

Table-N-E-7

Exp.	Syn	SM1	SM2	Rz	Ry	Y	AR	LCMS		
								method	RTime	Mass
N-e-187	NB1	Int.n-73	BRA1	2EtOPh	H	Et	2-Nap	C		441 (M <sup>+</sup> +1)
N-e-188	NA	N-e-187		2EtOPh	H	H	2-Nap	C		413 (M <sup>+</sup> +1)
N-e-189	NB1	Int.n-73	BRA3	2EtOPh	H	Et	1Me-5-Ind	C		444 (M <sup>+</sup> +1)
N-e-190	NA	N-e-189		2EtOPh	H	H	1Me-5-Ind	C		416 (M <sup>+</sup> +1)
N-e-191	NB1	Int.n-73	BRA6	2EtOPh	H	Et	1Me-5-1Hldz	C		445 (M <sup>+</sup> +1)
N-e-192	NA	N-e-191		2EtOPh	H	H	1Me-5-1Hldz	C		417 (M <sup>+</sup> +1)
N-e-193	NB1	Int.n-74	BRA1	3iPrOPh	H	Et	2-Nap	C		455 (M <sup>+</sup> +1)
N-e-194	NA	N-e-193		3iPrOPh	H	H	2-Nap	C		427 (M <sup>+</sup> +1)
N-e-195	NB1	Int.n-74	BRA2	3iPrOPh	H	Et	5-Ind	C		444 (M <sup>+</sup> +1)
N-e-196	NA	N-e-195		3iPrOPh	H	H	5-Ind	C		416 (M <sup>+</sup> +1)
N-e-197	NB1	Int.n-74	BRA3	3iPrOPh	H	Et	1Me-5-Ind	C		458 (M <sup>+</sup> +1)
N-e-198	NA	N-b-197		3iPrOPh	H	H	1Me-5-Ind	C		430 (M <sup>+</sup> +1)
N-e-199	NB1	Int.n-75	BRA3	3,5DFPh	H	Et	1Me-5-Ind	C		436 (M <sup>+</sup> +1)
N-e-200	NA	N-b-199		3,5DFPh	H	H	1Me-5-Ind	C		408 (M <sup>+</sup> +1)
N-e-201	NB1	Int.n-75	BRA5	3,5DFPh	H	Et	5-1Hldz	C		423 (M <sup>+</sup> +1)
N-e-202	NA	N-b-201		3,5DFPh	H	H	5-1Hldz	C		395 (M <sup>+</sup> +1)
N-e-203	NB1	Int.n-75	BRA6	3,5DFPh	H	Et	1Me-5-1Hldz	C		437 (M <sup>+</sup> +1)
N-e-204	NA	N-b-203		3,5DFPh	H	H	1Me-5-1Hldz	C		409 (M <sup>+</sup> +1)

## [Example N-f-1]

Synthesis of methyl 3-[3-(naphthalen-2-yl)-4-(N-phenylamino)phenyl]propionate  
(Compound No. N-f-1) (Synthesis method NB2)

A solution of Intermediate n-7 (306.1 mg) in dehydrated toluene (1 ml) was added with aniline (1 ml, TCI), palladium acetate (20.2 mg, WAKO), 2-(di-*t*-butylphosphine)biphenyl (39 mg, Across) and cesium carbonate (863.4 mg, WAKO), and stirred at 90°C for 18 hours. The reaction mixture was added with ethyl acetate (40 ml), and washed successively with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride and saturated brine. The organic layer was dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 4:1) to obtain the title compound (Compound No. N-f-1, 101.4 mg).

## [Examples N-f-1 to N-f-92]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of

the methods described in the present specification including the examples described above are shown in Table-N-F-1 and Table-N-F-2. In the tables, the compound numbers are mentioned in the columns indicated as "Exp.". In the tables, corresponding methods among the aforementioned synthesis methods are shown in the columns of "Syn" with symbols, the starting compounds 1 are mentioned in the columns of "SM1", and the starting compounds 2 are mentioned in the columns of "SM2".

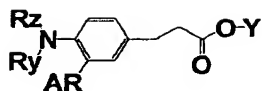


Table-N-F-1

Exp.	Syn	SM1	SM2	Rz	Ry	Y	AR	LCMS		
								method	RTime	Mass
N-f-1	NB2	Int.n-7	BRA14	Ph	H	Me	2-Nap	C		383 (M <sup>+</sup> +1)
N-f-2	NA	N-f-1		Ph	H	H	2-Nap	C		369 (M <sup>+</sup> +1)
N-f-3	NB2	Int.n-8	BRA14	Ph	H	Me	5-Ind	C		372 (M <sup>+</sup> +1)
N-f-4	NA	N-f-3		Ph	H	H	5-Ind	C		358 (M <sup>+</sup> +1)
N-f-5	NB2	Int.n-9	BRA14	Ph	H	Me	1Me-5-Ind	C		386 (M <sup>+</sup> +1)
N-f-6	NA	N-f-5		Ph	H	H	1Me-5-Ind	C		372 (M <sup>+</sup> +1)
N-f-7	NB2	Int.n-10	BRA14	Ph	H	Me	5-1HIdz	C		373 (M <sup>+</sup> +1)
N-f-8	NA	N-f-7		Ph	H	H	5-1HIdz	C		359 (M <sup>+</sup> +1)
N-f-9	NB2	Int.n-11	BRA14	Ph	H	Me	1Me-5-1HIdz	C		387 (M <sup>+</sup> +1)
N-f-10	NA	N-f-9		Ph	H	H	1Me-5-1HIdz	C		373 (M <sup>+</sup> +1)
N-f-11	NB2	N-f-1	CHO1	Ph	Me	Me	2-Nap	C		397 (M <sup>+</sup> +1)
N-f-12	NA	N-f-11		Ph	Me	H	2-Nap	C		383 (M <sup>+</sup> +1)
N-f-13	NB2	N-f-3	CHO1	Ph	Me	Me	1Me-5-Ind	C		400 (M <sup>+</sup> +1)
N-f-14	NA	N-f-13		Ph	Me	H	1Me-5-Ind	C		386 (M <sup>+</sup> +1)
N-f-15	NB2	N-f-5	CHO1	Ph	Me	Me	1Me-5-1HIdz	C		401 (M <sup>+</sup> +1)
N-f-16	NA	N-f-15		Ph	Me	H	1Me-5-1HIdz	C		387 (M <sup>+</sup> +1)
N-f-17	NB2	Int.n-7	BRA29	4MePh	H	Me	2-Nap	C		397 (M <sup>+</sup> +1)
N-f-18	NA	N-f-17		4MePh	H	H	2-Nap	C		383 (M <sup>+</sup> +1)
N-f-19	NB2	Int.n-9	BRA29	4MePh	H	Me	1Me-5-Ind	C		400 (M <sup>+</sup> +1)
N-f-20	NA	N-f-19		4MePh	H	H	1Me-5-Ind	C		386 (M <sup>+</sup> +1)
N-f-21	NB2	Int.n-11	BRA29	4MePh	H	Me	1Me-5-1HIdz	C		401 (M <sup>+</sup> +1)
N-f-22	NA	N-f-21		4MePh	H	H	1Me-5-1HIdz	C		387 (M <sup>+</sup> +1)
N-f-23	NB2	Int.n-7	BRA60	3MePh	H	Me	2-Nap	C		397 (M <sup>+</sup> +1)
N-f-24	NA	N-f-23		3MePh	H	H	2-Nap	C		383 (M <sup>+</sup> +1)
N-f-25	NB2	Int.n-9	BRA60	3MePh	H	Me	1Me-5-Ind	C		400 (M <sup>+</sup> +1)
N-f-26	NA	N-f-25		3MePh	H	H	1Me-5-Ind	C		386 (M <sup>+</sup> +1)
N-f-27	NB2	Int.n-11	BRA60	3MePh	H	Me	1Me-5-1HIdz	C		401 (M <sup>+</sup> +1)
N-f-28	NA	N-f-27		3MePh	H	H	1Me-5-1HIdz	C		387 (M <sup>+</sup> +1)
N-f-29	NB2	Int.n-7	BRA59	2MePh	H	Me	2-Nap	C		397 (M <sup>+</sup> +1)
N-f-30	NA	N-f-29		2MePh	H	H	2-Nap	C		383 (M <sup>+</sup> +1)
N-f-31	NB2	Int.n-8	BRA59	2MePh	H	Me	5-Ind	C		386 (M <sup>+</sup> +1)
N-f-32	NA	N-f-31		2MePh	H	H	5-Ind	C		372 (M <sup>+</sup> +1)
N-f-33	NB2	Int.n-10	BRA59	2MePh	H	Me	5-1HIdz	C		387 (M <sup>+</sup> +1)
N-f-34	NA	N-f-33		2MePh	H	H	5-1HIdz	C		373 (M <sup>+</sup> +1)
N-f-35	NB2	Int.n-7	BRA22	4FPh	H	Me	2-Nap	C		401 (M <sup>+</sup> +1)
N-f-36	NA	N-f-35		4FPh	H	H	2-Nap	C		387 (M <sup>+</sup> +1)
N-f-37	NB2	Int.n-8	BRA22	4FPh	H	Me	5-Ind	C		390 (M <sup>+</sup> +1)
N-f-38	NA	N-f-37		4FPh	H	H	5-Ind	C		376 (M <sup>+</sup> +1)
N-f-39	NB2	Int.n-9	BRA22	4FPh	H	Me	1Me-5-Ind	C		404 (M <sup>+</sup> +1)
N-f-40	NA	N-f-39		4FPh	H	H	1Me-5-Ind	C		390 (M <sup>+</sup> +1)
N-f-41	NB2	Int.n-7	BRA33	3FPh	H	Me	2-Nap	C		401 (M <sup>+</sup> +1)
N-f-42	NA	N-f-41		3FPh	H	H	2-Nap	C		387 (M <sup>+</sup> +1)
N-f-43	NB2	Int.n-10	BRA33	3FPh	H	Me	5-1HIdz	C		391 (M <sup>+</sup> +1)
N-f-44	NA	N-f-43		3FPh	H	H	5-1HIdz	C		377 (M <sup>+</sup> +1)
N-f-45	NB2	Int.n-11	BRA33	3FPh	H	Me	1Me-5-1HIdz	C		405 (M <sup>+</sup> +1)
N-f-46	NA	N-f-45		3FPh	H	H	1Me-5-1HIdz	C		391 (M <sup>+</sup> +1)



Table-N-F-2

Exp.	Syn	SM1	SM2	Rz	Ry	Y	AR	LCMS		
								method	RTime	Mass
N-f-47	NB2	Int.n-7	BRA32	2FPh	H	Me	2-Nap	C		401 (M <sup>+</sup> +1)
N-f-48	NA	N-f-47		2FPh	H	H	2-Nap	C		387 (M <sup>+</sup> +1)
N-f-49	NB2	Int.n-8	BRA32	2FPh	H	Me	5-Ind	C		390 (M <sup>+</sup> +1)
N-f-50	NA	N-f-49		2FPh	H	H	5-Ind	C		376 (M <sup>+</sup> +1)
N-f-51	NB2	Int.n-11	BRA32	2FPh	H	Me	1Me-5-1HIdz	C		405 (M <sup>+</sup> +1)
N-f-52	NA	N-f-51		2FPh	H	H	1Me-5-1HIdz	C		391 (M <sup>+</sup> +1)
N-f-53	NB2	Int.n-8	BRA19	4MeOPh	H	Me	5-Ind	C		402 (M <sup>+</sup> +1)
N-f-54	NA	N-f-53		4MeOPh	H	H	5-Ind	C		388 (M <sup>+</sup> +1)
N-f-55	NB2	Int.n-10	BRA19	4MeOPh	H	Me	5-1HIdz	C		403 (M <sup>+</sup> +1)
N-f-56	NA	N-f-55		4MeOPh	H	H	5-1HIdz	C		389 (M <sup>+</sup> +1)
N-f-57	NB2	Int.n-11	BRA19	4MeOPh	Me	Me	1Me-5-1HIdz	C		417 (M <sup>+</sup> +1)
N-f-58	NA	N-f-57		4MeOPh	Me	H	1Me-5-1HIdz	C		403 (M <sup>+</sup> +1)
N-f-59	NB2	Int.n-9	BRA37	3MeOPh	Me	Me	1Me-5-Ind	C		416 (M <sup>+</sup> +1)
N-f-60	NA	N-f-59		3MeOPh	Me	H	1Me-5-Ind	C		402 (M <sup>+</sup> +1)
N-f-61	NB2	Int.n-10	BRA37	3MeOPh	Me	Me	5-1HIdz	C		403 (M <sup>+</sup> +1)
N-f-62	NA	N-f-61		3MeOPh	Me	H	5-1HIdz	C		389 (M <sup>+</sup> +1)
N-f-63	NB2	Int.n-11	BRA37	3MeOPh	H	Me	1Me-5-1HIdz	C		417 (M <sup>+</sup> +1)
N-f-64	NA	N-f-63		3MeOPh	H	H	1Me-5-1HIdz	C		403 (M <sup>+</sup> +1)
N-f-65	NB2	Int.n-7	BRA38	2MeOPh	H	Me	2-Nap	C		413 (M <sup>+</sup> +1)
N-f-66	NA	N-f-65		2MeOPh	H	H	2-Nap	C		399 (M <sup>+</sup> +1)
N-f-67	NB2	Int.n-8	BRA38	2MeOPh	H	Me	5-Ind	C		402 (M <sup>+</sup> +1)
N-f-68	NA	N-f-67		2MeOPh	H	H	5-Ind	C		388 (M <sup>+</sup> +1)
N-f-69	NB2	Int.n-11	BRA38	2MeOPh	H	Me	1Me-5-1HIdz	C		417 (M <sup>+</sup> +1)
N-f-70	NA	N-f-69		2MeOPh	H	H	1Me-5-1HIdz	C		403 (M <sup>+</sup> +1)
N-f-71	NB2	Int.n-7	BRA41	4CF3Ph	H	Me	2-Nap	C		451 (M <sup>+</sup> +1)
N-f-72	NA	N-f-71		4CF3Ph	H	H	2-Nap	C		437 (M <sup>+</sup> +1)
N-f-73	NB2	Int.n-9	BRA41	4CF3Ph	H	Me	1Me-5-Ind	C		454 (M <sup>+</sup> +1)
N-f-74	NA	N-f-73		4CF3Ph	H	H	1Me-5-Ind	C		440 (M <sup>+</sup> +1)
N-f-75	NB2	Int.n-11	BRA41	4CF3Ph	H	Me	1Me-5-1HIdz	C		455 (M <sup>+</sup> +1)
N-f-76	NA	N-f-75		4CF3Ph	H	H	1Me-5-1HIdz	C		441 (M <sup>+</sup> +1)
N-f-77	NB2	Int.n-8	BRA88	4PhOPh	H	Me	5-Ind	C		464 (M <sup>+</sup> +1)
N-f-78	NA	N-f-77		4PhOPh	H	H	5-Ind	C		450 (M <sup>+</sup> +1)
N-f-79	NB2	Int.n-9	BRA88	4PhOPh	H	Me	1Me-5-Ind	C		478 (M <sup>+</sup> +1)
N-f-80	NA	N-f-79		4PhOPh	H	H	1Me-5-Ind	C		464 (M <sup>+</sup> +1)
N-f-81	NB2	Int.n-10	BRA88	4PhOPh	H	Me	5-1HIdz	C		465 (M <sup>+</sup> +1)
N-f-82	NA	N-f-81		4PhOPh	H	H	5-1HIdz	C		451 (M <sup>+</sup> +1)
N-f-83	NB2	Int.n-7	BRA61	2ClPh	H	Me	2-Nap	C		417 (M <sup>+</sup> +1)
N-f-84	NA	N-f-83		2ClPh	H	H	2-Nap	C		403 (M <sup>+</sup> +1)
N-f-85	NB2	Int.n-9	BRA61	2ClPh	H	Me	1Me-5-Ind	C		420 (M <sup>+</sup> +1)
N-f-86	NA	N-f-85		2ClPh	H	H	1Me-5-Ind	C		406 (M <sup>+</sup> +1)
N-f-87	NB2	Int.n-10	BRA61	2ClPh	H	Me	5-1HIdz	C		407 (M <sup>+</sup> +1)
N-f-88	NA	N-f-87		2ClPh	H	H	5-1HIdz	C		393 (M <sup>+</sup> +1)
N-f-89	NB2	Int.n-7	BRA73	3,5DMePh	H	Me	2-Nap	C		411 (M <sup>+</sup> +1)
N-f-90	NA	N-f-89		3,5DMePh	H	H	2-Nap	C		397 (M <sup>+</sup> +1)
N-f-91	NB2	Int.n-9	BRA73	3,5DMePh	H	Me	1Me-5-Ind	C		414 (M <sup>+</sup> +1)
N-f-92	NA	N-f-91		3,5DMePh	H	H	1Me-5-Ind	C		400 (M <sup>+</sup> +1)

[Example N-g-33]

Synthesis of methyl 3-[4-cyclopentylamino-3-methyl-5-(naphthalen-2-yl)phenyl]propionate (Compound No. N-e-33) (Synthesis method NB1)

According to the procedure described in the synthesis method of the compound of Example N-a-1 (Synthesis method NB) provided that the reaction was carried out for 18 hours, and the column chromatography was performed with hexane:ethyl acetate = 4:1, the compound of Example N-g-1 (91.6 mg), methyl boronate (140.0 mg, Ald), 2 M aqueous sodium carbonate (300  $\mu$ l) and (Ph<sub>3</sub>P)<sub>4</sub>Pd (75.5 mg) were reacted and treated to obtain the title compound (Compound No. N-g-33, 41.3 mg).

[Example N-g-251]

Synthesis of methyl 3-[4-(N-methyl-N-cyclopentylamino)-3-(N-methylamino)-5-(naphthalen-2-yl)phenyl]propionate (Compound No. N-g-251) (Synthesis method NN1)

A solution of Compound No. N-g-131 (102 mg) in DMF (3 ml) was added with 60% sodium hydride (7 mg) under ice cooling, and stirred for 10 minutes. This reaction mixture was added with methyl iodide (17  $\mu$ l), stirred for 10 minutes, then warmed to room temperature, and further stirred for 2 hours. The reaction mixture was poured into water, and added with ethyl acetate (30 ml) for extraction. The organic layer was washed successively with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride and saturated brine, and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 3:1) to obtain the title compound (Compound No. N-g-251, 30 mg).

[Example N-g-285]

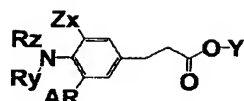
Synthesis of methyl 3-[3-(N-dimethylamino)-4-(N-methyl-N-cyclopentylamino)-5-(naphthalen-2-yl)phenyl]propionate (Compound No. N-g-285) (Synthesis method

NN2)

A solution of Compound No. N-g-131 (102 mg) in DMF (3 ml) was added with 60% sodium hydride (20 mg) under ice cooling, and stirred for 10 minutes. This reaction mixture was added dropwise with methyl iodide (100  $\mu$ l), stirred for 10 minutes, then warmed to room temperature, and further stirred for 16 hours. The reaction mixture was poured into water, and added with ethyl acetate (30 ml) for extraction. The organic layer was washed successively with saturated aqueous sodium hydrogencarbonate, saturated aqueous ammonium chloride and saturated brine, and dried, and then the solvent was evaporated under reduced pressure. The residue was purified by column chromatography (Quad, hexane:ethyl acetate = 3:1) to obtain the title compound (Compound No. N-g-285, 80 mg).

[Examples N-g-1 to N-g-318]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification including the examples described above are shown in Table N-G-1 to Table N-G-7. In the tables, the compound numbers are mentioned in the columns indicated as "Exp.". In the tables, used methods among the aforementioned synthesis methods are shown in the columns of "Syn" with symbols, the starting compounds 1 are mentioned in the columns of "SM1", and the starting compounds 2 are mentioned in the columns of "SM2".



Tablg-N-G-1

Exp.	Syn	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									method	RTime	Mass
N-g-1	NB1	Int.n-39	BRA1	cPen	H	Me	Br	2-Nap	C		452 (M <sup>+</sup> )
N-g-2	NA	N-g-1		cPen	H	H	Br	2-Nap	C		438 (M <sup>+</sup> )
N-g-3	NB1	Int.n-39	BRA2	cPen	H	Me	Br	5-Ind	C		441 (M <sup>+</sup> )
N-g-4	NA	N-g-3		cPen	H	H	Br	5-Ind	C		427 (M <sup>+</sup> )
N-g-5	NB1	Int.n-39	BRA3	cPen	H	Me	Br	1Me-5-Ind	C		455 (M <sup>+</sup> )
N-g-6	NA	N-g-5		cPen	H	H	Br	1Me-5-Ind	C		441 (M <sup>+</sup> )
N-g-7	NB1	Int.n-39	BRA5	cPen	H	Me	Br	5-1HIdz	C		442 (M <sup>+</sup> )
N-g-8	NA	N-g-7		cPen	H	H	Br	5-1HIdz	C		428 (M <sup>+</sup> )
N-g-9	NB1	Int.n-39	BRA6	cPen	H	Me	Br	1Me-5-1HIdz	C		456 (M <sup>+</sup> )
N-g-10	NA	N-g-9		cPen	H	H	Br	1Me-5-1HIdz	C		442 (M <sup>+</sup> )
N-g-11	NB1	Int.n-39	BRA11	cPen	H	Me	Br	6-Qu	C		453 (M <sup>+</sup> )
N-g-12	NA	N-g-11		cPen	H	H	Br	6-Qu	C		439 (M <sup>+</sup> )
N-g-13	NC2	N-g-1	CHO1	cPen	Me	Me	Br	2-Nap	C		466 (M <sup>+</sup> )
N-g-14	NA	N-g-13		cPen	Me	H	Br	2-Nap	C		452 (M <sup>+</sup> )
N-g-15	NC2	N-g-5	CHO1	cPen	Me	Me	Br	1Me-5-Ind	C		455 (M <sup>+</sup> )
N-g-16	NA	N-g-15		cPen	Me	H	Br	1Me-5-Ind	C		441 (M <sup>+</sup> )
N-g-17	NB1	Int.n-41	BRA2	nPr	H	Me	Br	5-Ind	C		415 (M <sup>+</sup> )
N-g-18	NA	N-g-17		nPr	H	H	Br	5-Ind	C		401 (M <sup>+</sup> )
N-g-19	NB1	Int.n-41	BRA3	nPr	H	Me	Br	1Me-5-Ind	C		429 (M <sup>+</sup> )
N-g-20	NA	N-g-19		nPr	H	H	Br	1Me-5-Ind	C		415 (M <sup>+</sup> )
N-g-21	NB1	Int.n-41	BRA5	nPr	H	Me	Br	5-1HIdz	C		416 (M <sup>+</sup> )
N-g-22	NA	N-g-21		nPr	H	H	Br	5-1HIdz	C		402 (M <sup>+</sup> )
N-g-23	NB1	Int.n-41	BRA11	nPr	H	Me	Br	6-Qu	C		427 (M <sup>+</sup> )
N-g-24	NA	N-g-23		nPr	H	H	Br	6-Qu	C		413 (M <sup>+</sup> )
N-g-25	NB1	Int.n-43	BRA1	iPr	H	Me	Br	2-Nap	C		426 (M <sup>+</sup> )
N-g-26	NA	N-g-25		iPr	H	H	Br	2-Nap	C		412 (M <sup>+</sup> )
N-g-27	NB1	Int.n-43	BRA2	iPr	H	Me	Br	5-Ind	C		415 (M <sup>+</sup> )
N-g-28	NA	N-g-27		iPr	H	H	Br	5-Ind	C		401 (M <sup>+</sup> )
N-g-29	NB1	Int.n-43	BRA6	iPr	H	Me	Br	1Me-5-1HIdz	C		430 (M <sup>+</sup> )
N-g-30	NA	N-g-29		iPr	H	H	Br	1Me-5-1HIdz	C		416 (M <sup>+</sup> )
N-g-31	NB1	Int.n-43	BRA10	iPr	H	Me	Br	3-Qu	C		427 (M <sup>+</sup> )
N-g-32	NA	N-g-31		iPr	H	H	Br	3-Qu	C		413 (M <sup>+</sup> )
N-g-33	NB1	N-g-1	BRA13	cPen	H	Me	Me	2-Nap	C		388 (M <sup>+</sup> +1)
N-g-34	NA	N-g-33		cPen	H	H	Me	2-Nap	C		374 (M <sup>+</sup> +1)
N-g-35	NB1	N-g-3	BRA13	cPen	H	Me	Me	5-Ind	C		377 (M <sup>+</sup> +1)
N-g-36	NA	N-g-35		cPen	H	H	Me	5-Ind	C		363 (M <sup>+</sup> +1)
N-g-37	NB1	N-g-5	BRA13	cPen	H	Me	Me	1Me-5-Ind	C		391 (M <sup>+</sup> +1)
N-g-38	NA	N-g-37		cPen	H	H	Me	1Me-5-Ind	C		377 (M <sup>+</sup> +1)
N-g-39	NB1	N-g-7	BRA13	cPen	H	Me	Me	5-1HIdz	C		378 (M <sup>+</sup> +1)
N-g-40	NA	N-g-39		cPen	H	H	Me	5-1HIdz	C		364 (M <sup>+</sup> +1)
N-g-41	NB1	N-g-9	BRA13	cPen	H	Me	Me	1Me-5-1HIdz	C		392 (M <sup>+</sup> +1)
N-g-42	NA	N-g-41		cPen	H	H	Me	1Me-5-1HIdz	C		378 (M <sup>+</sup> +1)
N-g-43	NC2	N-g-37	CHO1	cPen	Me	Me	Me	1Me-5-Ind	C		405 (M <sup>+</sup> +1)
N-g-44	NA	N-g-43		cPen	Me	H	Me	1Me-5-Ind	C		391 (M <sup>+</sup> +1)

Tablg-N-G-2

Exp.	Syn	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									method	RTime	Mass
N-g-45	NB1	Intn-77	BRA1	cPen	H	Me	NO2	2-Nap	C		419 (M <sup>+</sup> +1)
N-g-46	NA	N-g-45		cPen	H	H	NO2	2-Nap	C		405 (M <sup>+</sup> +1)
N-g-47	NB1	Intn-77	BRA2	cPen	H	Me	NO2	5-Ind	C		408 (M <sup>+</sup> +1)
N-g-48	NA	N-g-47		cPen	H	H	NO2	5-Ind	C		394 (M <sup>+</sup> +1)
N-g-49	NB1	Intn-77	BRA3	cPen	H	Me	NO2	1Me-5-Ind	C		422 (M <sup>+</sup> +1)
N-g-50	NA	N-g-49		cPen	H	H	NO2	1Me-5-Ind	C		408 (M <sup>+</sup> +1)
N-g-51	NB1	Intn-77	BRA5	cPen	H	Me	NO2	5-1HIdz	C		409 (M <sup>+</sup> +1)
N-g-52	NA	N-g-51		cPen	H	H	NO2	5-1HIdz	C		395 (M <sup>+</sup> +1)
N-g-53	NB1	Intn-77	BRA6	cPen	H	Me	NO2	1Me-5-1HIdz	C		423 (M <sup>+</sup> +1)
N-g-54	NA	N-g-53		cPen	H	H	NO2	1Me-5-1HIdz	C		409 (M <sup>+</sup> +1)
N-g-55	NB1	Intn-78	BRA1	nPr	H	Me	NO2	2-Nap	C		393 (M <sup>+</sup> +1)
N-g-56	NA	N-g-55		nPr	H	H	NO2	2-Nap	C		379 (M <sup>+</sup> +1)
N-g-57	NB1	Intn-78	BRA2	nPr	H	Me	NO2	5-Ind	C		382 (M <sup>+</sup> +1)
N-g-58	NA	N-g-57		nPr	H	H	NO2	5-Ind	C		368 (M <sup>+</sup> +1)
N-g-59	NB1	Intn-78	BRA3	nPr	H	Me	NO2	1Me-5-Ind	C		396 (M <sup>+</sup> +1)
N-g-60	NA	N-g-59		nPr	H	H	NO2	1Me-5-Ind	C		382 (M <sup>+</sup> +1)
N-g-61	NB1	Intn-78	BRA5	nPr	H	Me	NO2	5-1HIdz	C		383 (M <sup>+</sup> +1)
N-g-62	NA	N-g-61		nPr	H	H	NO2	5-1HIdz	C		369 (M <sup>+</sup> +1)
N-g-63	NB1	Intn-78	BRA6	nPr	H	Me	NO2	1Me-5-1HIdz	C		397 (M <sup>+</sup> +1)
N-g-64	NA	N-g-63		nPr	H	H	NO2	1Me-5-1HIdz	C		383 (M <sup>+</sup> +1)
N-g-65	NB1	Intn-79	BRA1	iPr	H	Me	NO2	2-Nap	C		393 (M <sup>+</sup> +1)
N-g-66	NA	N-g-65		iPr	H	H	NO2	2-Nap	C		379 (M <sup>+</sup> +1)
N-g-67	NB1	Intn-79	BRA2	iPr	H	Me	NO2	5-Ind	C		382 (M <sup>+</sup> +1)
N-g-68	NA	N-g-67		iPr	H	H	NO2	5-Ind	C		368 (M <sup>+</sup> +1)
N-g-69	NB1	Intn-79	BRA3	iPr	H	Me	NO2	1Me-5-Ind	C		396 (M <sup>+</sup> +1)
N-g-70	NA	N-g-69		iPr	H	H	NO2	1Me-5-Ind	C		382 (M <sup>+</sup> +1)
N-g-71	NB1	Intn-79	BRA5	iPr	H	Me	NO2	5-1HIdz	C		383 (M <sup>+</sup> +1)
N-g-72	NA	N-g-71		iPr	H	H	NO2	5-1HIdz	C		369 (M <sup>+</sup> +1)
N-g-73	NB1	Intn-79	BRA6	iPr	H	Me	NO2	1Me-5-1HIdz	C		397 (M <sup>+</sup> +1)
N-g-74	NA	N-g-73		iPr	H	H	NO2	1Me-5-1HIdz	C		383 (M <sup>+</sup> +1)
N-g-75	NB1	Intn-83	BRA1	cPen	Me	Me	NO2	2-Nap	C		433 (M <sup>+</sup> +1)
N-g-76	NA	N-g-75		cPen	Me	H	NO2	2-Nap	C		419 (M <sup>+</sup> +1)
N-g-77	NB1	Intn-83	BRA3	cPen	Me	Me	NO2	1Me-5-Ind	C		436 (M <sup>+</sup> +1)
N-g-78	NA	N-g-77		cPen	Me	H	NO2	1Me-5-Ind	C		422 (M <sup>+</sup> +1)
N-g-79	NB1	Intn-83	BRA6	cPen	Me	Me	NO2	1Me-5-1HIdz	C		437 (M <sup>+</sup> +1)
N-g-80	NA	N-g-79		cPen	Me	H	NO2	1Me-5-1HIdz	C		423 (M <sup>+</sup> +1)
N-g-81	NB1	Intn-84	BRA1	nPr	Me	Me	NO2	2-Nap	C		407 (M <sup>+</sup> +1)
N-g-82	NA	N-g-81		nPr	Me	H	NO2	2-Nap	C		393 (M <sup>+</sup> +1)
N-g-83	NB1	Intn-84	BRA2	nPr	Me	Me	NO2	5-Ind	C		396 (M <sup>+</sup> +1)
N-g-84	NA	N-g-83		nPr	Me	H	NO2	5-Ind	C		382 (M <sup>+</sup> +1)
N-g-85	NB1	Intn-84	BRA3	nPr	Me	Me	NO2	1Me-5-Ind	C		410 (M <sup>+</sup> +1)
N-g-86	NA	N-g-85		nPr	Me	H	NO2	1Me-5-Ind	C		396 (M <sup>+</sup> +1)
N-g-87	NB1	Intn-84	BRA5	nPr	Me	Me	NO2	5-1HIdz	C		397 (M <sup>+</sup> +1)
N-g-88	NA	N-g-87		nPr	Me	H	NO2	5-1HIdz	C		383 (M <sup>+</sup> +1)
N-g-89	NB1	Intn-84	BRA6	nPr	Me	Me	NO2	1Me-5-1HIdz	C		411 (M <sup>+</sup> +1)
N-g-90	NA	N-g-89		nPr	Me	H	NO2	1Me-5-1HIdz	C		397 (M <sup>+</sup> +1)
N-g-91	NB1	Intn-85	BRA1	iPr	Me	Me	NO2	2-Nap	C		407 (M <sup>+</sup> +1)
N-g-92	NA	N-g-91		iPr	Me	H	NO2	2-Nap	C		393 (M <sup>+</sup> +1)

Tabl-N-G-3

Exp.	Syn	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									method	RTime	Mass
N-g-93	NB1	Int.n-85	BRA2	iPr	Me	Me	NO2	5-Ind	C		396 (M <sup>+</sup> +1)
N-g-94	NA	N-g-93		iPr	Me	H	NO2	5-Ind	C		382 (M <sup>+</sup> +1)
N-g-95	NB1	Int.n-85	BRA3	iPr	Me	Me	NO2	1Me-5-Ind	C		410 (M <sup>+</sup> +1)
N-g-96	NA	N-g-95		iPr	Me	H	NO2	1Me-5-Ind	C		396 (M <sup>+</sup> +1)
N-g-97	NB1	Int.n-85	BRA5	iPr	Me	Me	NO2	5-1HIdz	C		397 (M <sup>+</sup> +1)
N-g-98	NA	N-g-97		iPr	Me	H	NO2	5-1HIdz	C		383 (M <sup>+</sup> +1)
N-g-99	NB1	Int.n-85	BRA6	iPr	Me	Me	NO2	1Me-5-1HIdz	C		411 (M <sup>+</sup> +1)
N-g-100	NA	N-g-99		iPr	Me	H	NO2	1Me-5-1HIdz	C		397 (M <sup>+</sup> +1)
N-g-101	ND1	N-g-45		cPen	H	Me	NH2	2-Nap	C		389 (M <sup>+</sup> +1)
N-g-102	NA	N-g-101		cPen	H	H	NH2	2-Nap	C		375 (M <sup>+</sup> +1)
N-g-103	ND1	N-g-47		cPen	H	Me	NH2	5-Ind	C		378 (M <sup>+</sup> +1)
N-g-104	NA	N-g-103		cPen	H	H	NH2	5-Ind	C		364 (M <sup>+</sup> +1)
N-g-105	ND1	N-g-49		cPen	H	Me	NH2	1Me-5-Ind	C		392 (M <sup>+</sup> +1)
N-g-106	NA	N-g-105		cPen	H	H	NH2	1Me-5-Ind	C		378 (M <sup>+</sup> +1)
N-g-107	ND1	N-g-51		cPen	H	Me	NH2	5-1HIdz	C		379 (M <sup>+</sup> +1)
N-g-108	NA	N-g-107		cPen	H	H	NH2	5-1HIdz	C		365 (M <sup>+</sup> +1)
N-g-109	ND1	N-g-53		cPen	H	Me	NH2	1Me-5-1HIdz	C		393 (M <sup>+</sup> +1)
N-g-110	NA	N-g-109		cPen	H	H	NH2	1Me-5-1HIdz	C		379 (M <sup>+</sup> +1)
N-g-111	ND1	N-g-55		nPr	H	Me	NH2	2-Nap	C		363 (M <sup>+</sup> +1)
N-g-112	NA	N-g-111		nPr	H	H	NH2	2-Nap	C		349 (M <sup>+</sup> +1)
N-g-113	ND1	N-g-57		nPr	H	Me	NH2	5-Ind	C		352 (M <sup>+</sup> +1)
N-g-114	NA	N-g-113		nPr	H	H	NH2	5-Ind	C		338 (M <sup>+</sup> +1)
N-g-115	ND1	N-g-59		nPr	H	Me	NH2	1Me-5-Ind	C		366 (M <sup>+</sup> +1)
N-g-116	NA	N-g-115		nPr	H	H	NH2	1Me-5-Ind	C		352 (M <sup>+</sup> +1)
N-g-117	ND1	N-g-61		nPr	H	Me	NH2	5-1HIdz	C		353 (M <sup>+</sup> +1)
N-g-118	NA	N-g-117		nPr	H	H	NH2	5-1HIdz	C		339 (M <sup>+</sup> +1)
N-g-119	ND1	N-g-63		nPr	H	Me	NH2	1Me-5-1HIdz	C		367 (M <sup>+</sup> +1)
N-g-120	NA	N-g-119		nPr	H	H	NH2	1Me-5-1HIdz	C		353 (M <sup>+</sup> +1)
N-g-121	ND1	N-g-65		iPr	H	Me	NH2	2-Nap	C		363 (M <sup>+</sup> +1)
N-g-122	NA	N-g-121		iPr	H	H	NH2	2-Nap	C		349 (M <sup>+</sup> +1)
N-g-123	ND1	N-g-67		iPr	H	Me	NH2	5-Ind	C		352 (M <sup>+</sup> +1)
N-g-124	NA	N-g-123		iPr	H	H	NH2	5-Ind	C		338 (M <sup>+</sup> +1)
N-g-125	ND1	N-g-69		iPr	H	Me	NH2	1Me-5-Ind	C		366 (M <sup>+</sup> +1)
N-g-126	NA	N-g-125		iPr	H	H	NH2	1Me-5-Ind	C		352 (M <sup>+</sup> +1)
N-g-127	ND1	N-g-71		iPr	H	Me	NH2	5-1HIdz	C		353 (M <sup>+</sup> +1)
N-g-128	NA	N-g-127		iPr	H	H	NH2	5-1HIdz	C		339 (M <sup>+</sup> +1)
N-g-129	ND1	N-g-73		iPr	H	Me	NH2	1Me-5-1HIdz	C		367 (M <sup>+</sup> +1)
N-g-130	NA	N-g-129		iPr	H	H	NH2	1Me-5-1HIdz	C		353 (M <sup>+</sup> +1)
N-g-131	NC1	N-g-75		cPen	Me	Me	NH2	2-Nap	C		389 (M <sup>+</sup> +1)
N-g-132	NA	N-g-131		cPen	Me	H	NH2	2-Nap	C		375 (M <sup>+</sup> +1)
N-g-133	NC1	N-g-77		cPen	Me	Me	NH2	1Me-5-Ind	C		392 (M <sup>+</sup> +1)
N-g-134	NA	N-g-133		cPen	Me	H	NH2	1Me-5-Ind	C		378 (M <sup>+</sup> +1)
N-g-135	NC1	N-g-79		cPen	Me	Me	NH2	5-1HIdz	C		379 (M <sup>+</sup> +1)
N-g-136	NA	N-g-135		cPen	Me	H	NH2	5-1HIdz	C		365 (M <sup>+</sup> +1)
N-g-137	NC1	N-g-81		cPen	Me	Me	NH2	1Me-5-1HIdz	C		393 (M <sup>+</sup> +1)
N-g-138	NA	N-g-137		cPen	Me	H	NH2	1Me-5-1HIdz	C		379 (M <sup>+</sup> +1)

Tablg-N-G-4

Exp.	Syn	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									method	RTime	Mass
N-g-139	ND1	N-g-83		nPr	Me	Me	NH2	2-Nap	C		363 (M <sup>+</sup> +1)
N-g-140	NA	N-g-139		nPr	Me	H	NH2	2-Nap	C		349 (M <sup>+</sup> +1)
N-g-141	ND1	N-g-85		nPr	Me	Me	NH2	5-Ind	C		352 (M <sup>+</sup> +1)
N-g-142	NA	N-g-141		nPr	Me	H	NH2	5-Ind	C		338 (M <sup>+</sup> +1)
N-g-143	ND1	N-g-87		nPr	Me	Me	NH2	1Me-5-Ind	C		366 (M <sup>+</sup> +1)
N-g-144	NA	N-g-143		nPr	Me	H	NH2	1Me-5-Ind	C		352 (M <sup>+</sup> +1)
N-g-145	ND1	N-g-89		nPr	Me	Me	NH2	5-1HIdz	C		353 (M <sup>+</sup> +1)
N-g-146	NA	N-g-145		nPr	Me	H	NH2	5-1HIdz	C		339 (M <sup>+</sup> +1)
N-g-147	ND1	N-g-91		nPr	Me	Me	NH2	1Me-5-1HIdz	C		367 (M <sup>+</sup> +1)
N-g-148	NA	N-g-147		nPr	Me	H	NH2	1Me-5-1HIdz	C		353 (M <sup>+</sup> +1)
N-g-149	ND1	N-g-93		iPr	Me	Me	NH2	2-Nap	C		363 (M <sup>+</sup> +1)
N-g-150	NA	N-g-149		iPr	Me	H	NH2	2-Nap	C		349 (M <sup>+</sup> +1)
N-g-151	ND1	N-g-95		iPr	Me	Me	NH2	1Me-5-Ind	C		366 (M <sup>+</sup> +1)
N-g-152	NA	N-g-151		iPr	Me	H	NH2	1Me-5-Ind	C		352 (M <sup>+</sup> +1)
N-g-153	ND1	N-g-97		iPr	Me	Me	NH2	5-1HIdz	C		353 (M <sup>+</sup> +1)
N-g-154	NA	N-g-153		iPr	Me	H	NH2	5-1HIdz	C		339 (M <sup>+</sup> +1)
N-g-155	ND1	N-g-99		iPr	Me	Me	NH2	1Me-5-1HIdz	C		367 (M <sup>+</sup> +1)
N-g-156	NA	N-g-155		iPr	Me	H	NH2	1Me-5-1HIdz	C		353 (M <sup>+</sup> +1)
N-g-157	NB1	Int.n-80	BRA1	2-Indane	H	Me	NO2	2-Nap	C		467 (M <sup>+</sup> +1)
N-g-158	NA	N-g-157		2-Indane	H	H	NO2	2-Nap	C		453 (M <sup>+</sup> +1)
N-g-159	NB1	Int.n-80	BRA2	2-Indane	H	Me	NO2	5-Ind	C		456 (M <sup>+</sup> +1)
N-g-160	NA	N-g-159		2-Indane	H	H	NO2	5-Ind	C		442 (M <sup>+</sup> +1)
N-g-161	NB1	Int.n-80	BRA3	2-Indane	H	Me	NO2	1Me-5-Ind	C		470 (M <sup>+</sup> +1)
N-g-162	NA	N-g-161		2-Indane	H	H	NO2	1Me-5-Ind	C		456 (M <sup>+</sup> +1)
N-g-163	NB1	Int.n-80	BRA5	2-Indane	H	Me	NO2	5-1HIdz	C		457 (M <sup>+</sup> +1)
N-g-164	NA	N-g-163		2-Indane	H	H	NO2	5-1HIdz	C		443 (M <sup>+</sup> +1)
N-g-165	NB1	Int.n-80	BRA6	2-Indane	H	Me	NO2	1Me-5-1HIdz	C		471 (M <sup>+</sup> +1)
N-g-166	NA	N-g-165		2-Indane	H	H	NO2	1Me-5-1HIdz	C		457 (M <sup>+</sup> +1)
N-g-167	NB1	Int.n-81	BRA2	cHex	H	Me	NO2	5-Ind	C		422 (M <sup>+</sup> +1)
N-g-168	NA	N-g-167		cHex	H	H	NO2	5-Ind	C		408 (M <sup>+</sup> +1)
N-g-169	NB1	Int.n-81	BRA3	cHex	H	Me	NO2	1Me-5-Ind	C		436 (M <sup>+</sup> +1)
N-g-170	NA	N-g-169		cHex	H	H	NO2	1Me-5-Ind	C		422 (M <sup>+</sup> +1)
N-g-171	NB1	Int.n-81	BRA5	cHex	H	Me	NO2	5-1HIdz	C		423 (M <sup>+</sup> +1)
N-g-172	NA	N-g-171		cHex	H	H	NO2	5-1HIdz	C		409 (M <sup>+</sup> +1)
N-g-173	NB1	Int.n-81	BRA6	cHex	H	Me	NO2	1Me-5-1HIdz	C		437 (M <sup>+</sup> +1)
N-g-174	NA	N-g-173		cHex	H	H	NO2	1Me-5-1HIdz	C		423 (M <sup>+</sup> +1)
N-g-175	NB1	Int.n-82	BRA1	2(Me)cHex	H	Me	NO2	2-Nap	C		447 (M <sup>+</sup> +1)
N-g-176	NA	N-g-175		2(Me)cHex	H	H	NO2	2-Nap	C		433 (M <sup>+</sup> +1)
N-g-177	NB1	Int.n-82	BRA2	2(Me)cHex	H	Me	NO2	5-Ind	C		436 (M <sup>+</sup> +1)
N-g-178	NA	N-g-177		2(Me)cHex	H	H	NO2	5-Ind	C		422 (M <sup>+</sup> +1)
N-g-179	NB1	Int.n-82	BRA3	2(Me)cHex	H	Me	NO2	1Me-5-Ind	C		450 (M <sup>+</sup> +1)
N-g-180	NA	N-g-179		2(Me)cHex	H	H	NO2	1Me-5-Ind	C		436 (M <sup>+</sup> +1)
N-g-181	NB1	Int.n-82	BRA5	2(Me)cHex	H	Me	NO2	5-1HIdz	C		437 (M <sup>+</sup> +1)
N-g-182	NA	N-g-181		2(Me)cHex	H	H	NO2	5-1HIdz	C		423 (M <sup>+</sup> +1)
N-g-183	NB1	Int.n-82	BRA6	2(Me)cHex	H	Me	NO2	1Me-5-1HIdz	C		451 (M <sup>+</sup> +1)
N-g-184	NA	N-g-183		2(Me)cHex	H	H	NO2	1Me-5-1HIdz	C		437 (M <sup>+</sup> +1)

Tablg-N-G-5

Exp.	Syn	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									method	RTime	Mass
N-g-185	NB1	Int.n-86	BRA1	2-Indane	Me	Me	NO2	2-Nap	C		481 (M <sup>+</sup> +1)
N-g-186	NA	N-g-185		2-Indane	Me	H	NO2	2-Nap	C		467 (M <sup>+</sup> +1)
N-g-187	NB1	Int.n-86	BRA3	2-Indane	Me	Me	NO2	1Me-5-Ind	C		484 (M <sup>+</sup> +1)
N-g-188	NA	N-g-187		2-Indane	Me	H	NO2	1Me-5-Ind	C		470 (M <sup>+</sup> +1)
N-g-189	NB1	Int.n-86	BRA5	2-Indane	Me	Me	NO2	5-1HIdz	C		471 (M <sup>+</sup> +1)
N-g-190	NA	N-g-189		2-Indane	Me	H	NO2	5-1HIdz	C		457 (M <sup>+</sup> +1)
N-g-191	NB1	Int.n-86	BRA6	2-Indane	Me	Me	NO2	1Me-5-1HIdz	C		485 (M <sup>+</sup> +1)
N-g-192	NA	N-g-191		2-Indane	Me	H	NO2	1Me-5-1HIdz	C		471 (M <sup>+</sup> +1)
N-g-193	NB1	Int.n-87	BRA1	cHex	Me	Me	NO2	2-Nap	C		447 (M <sup>+</sup> +1)
N-g-194	NA	N-g-193		cHex	Me	H	NO2	2-Nap	C		433 (M <sup>+</sup> +1)
N-g-195	NB1	Int.n-87	BRA3	cHex	Me	Me	NO2	1Me-5-Ind	C		450 (M <sup>+</sup> +1)
N-g-196	NA	N-g-195		cHex	Me	H	NO2	1Me-5-Ind	C		436 (M <sup>+</sup> +1)
N-g-197	NB1	Int.n-87	BRA5	cHex	Me	Me	NO2	5-1HIdz	C		437 (M <sup>+</sup> +1)
N-g-198	NA	N-g-197		cHex	Me	H	NO2	5-1HIdz	C		423 (M <sup>+</sup> +1)
N-g-199	NB1	Int.n-87	BRA6	cHex	Me	Me	NO2	1Me-5-1HIdz	C		451 (M <sup>+</sup> +1)
N-g-200	NA	N-g-199		cHex	Me	H	NO2	1Me-5-1HIdz	C		437 (M <sup>+</sup> +1)
N-g-201	NB1	Int.n-88	BRA1	4(Me)cHex	Me	Me	NO2	2-Nap	C		461 (M <sup>+</sup> +1)
N-g-202	NA	N-g-201		4(Me)cHex	Me	H	NO2	2-Nap	C		447 (M <sup>+</sup> +1)
N-g-203	NB1	Int.n-88	BRA3	4(Me)cHex	Me	Me	NO2	1Me-5-Ind	C		464 (M <sup>+</sup> +1)
N-g-204	NA	N-g-203		4(Me)cHex	Me	H	NO2	1Me-5-Ind	C		450 (M <sup>+</sup> +1)
N-g-205	NB1	Int.n-88	BRA6	4(Me)cHex	Me	Me	NO2	1Me-5-1HIdz	C		465 (M <sup>+</sup> +1)
N-g-206	NA	N-g-205		4(Me)cHex	Me	H	NO2	1Me-5-1HIdz	C		451 (M <sup>+</sup> +1)
N-g-207	ND1	N-g-159		2-Indane	H	Me	NH2	5-Ind	C		426 (M <sup>+</sup> +1)
N-g-208	NA	N-g-207		2-Indane	H	H	NH2	5-Ind	C		412 (M <sup>+</sup> +1)
N-g-209	ND1	N-g-161		2-Indane	H	Me	NH2	1Me-5-Ind	C		440 (M <sup>+</sup> +1)
N-g-210	NA	N-g-209		2-Indane	H	H	NH2	1Me-5-Ind	C		426 (M <sup>+</sup> +1)
N-g-211	ND1	N-g-163		2-Indane	H	Me	NH2	5-1HIdz	C		427 (M <sup>+</sup> +1)
N-g-212	NA	N-g-211		2-Indane	H	H	NH2	5-1HIdz	C		413 (M <sup>+</sup> +1)
N-g-213	ND1	N-g-165		2-Indane	H	Me	NH2	1Me-5-1HIdz	C		441 (M <sup>+</sup> +1)
N-g-214	NA	N-g-213		2-Indane	H	H	NH2	1Me-5-1HIdz	C		427 (M <sup>+</sup> +1)
N-g-215	ND1	N-g-167		cHex	H	Me	NH2	5-Ind	C		392 (M <sup>+</sup> +1)
N-g-216	NA	N-g-215		cHex	H	H	NH2	5-Ind	C		378 (M <sup>+</sup> +1)
N-g-217	ND1	N-g-169		cHex	H	Me	NH2	1Me-5-Ind	C		406 (M <sup>+</sup> +1)
N-g-218	NA	N-g-217		cHex	H	H	NH2	1Me-5-Ind	C		392 (M <sup>+</sup> +1)
N-g-219	ND1	N-g-171		cHex	H	Me	NH2	5-1HIdz	C		393 (M <sup>+</sup> +1)
N-g-220	NA	N-g-219		cHex	H	H	NH2	5-1HIdz	C		379 (M <sup>+</sup> +1)
N-g-221	ND1	N-g-173		cHex	H	Me	NH2	1Me-5-1HIdz	C		407 (M <sup>+</sup> +1)
N-g-222	NA	N-g-221		cHex	H	H	NH2	1Me-5-1HIdz	C		393 (M <sup>+</sup> +1)
N-g-223	ND1	N-g-177		4(Me)cHex	H	Me	NH2	5-Ind	C		406 (M <sup>+</sup> +1)
N-g-224	NA	N-g-223		4(Me)cHex	H	H	NH2	5-Ind	C		392 (M <sup>+</sup> +1)
N-g-225	ND1	N-g-179		4(Me)cHex	H	Me	NH2	1Me-5-Ind	C		420 (M <sup>+</sup> +1)
N-g-226	NA	N-g-225		4(Me)cHex	H	H	NH2	1Me-5-Ind	C		406 (M <sup>+</sup> +1)
N-g-227	ND1	N-g-181		4(Me)cHex	H	Me	NH2	5-1HIdz	C		407 (M <sup>+</sup> +1)
N-g-228	NA	N-g-227		4(Me)cHex	H	H	NH2	5-1HIdz	C		393 (M <sup>+</sup> +1)
N-g-229	ND1	N-g-183		4(Me)cHex	H	Me	NH2	1Me-5-1HIdz	C		421 (M <sup>+</sup> +1)
N-g-230	NA	N-g-229		4(Me)cHex	H	H	NH2	1Me-5-1HIdz	C		407 (M <sup>+</sup> +1)



Table-N-G-6

Exp.	Syn	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									method	RTime	Mass
N-g-231	ND1	N-g-185		2-Indane	Me	Me	NH2	2-Nap	C		451 (M <sup>+</sup> +1)
N-g-232	NA	N-g-231		2-Indane	Me	H	NH2	2-Nap	C		437 (M <sup>+</sup> +1)
N-g-233	ND1	N-g-187		2-Indane	Me	Me	NH2	1Me-5-Ind	C		454 (M <sup>+</sup> +1)
N-g-234	NA	N-g-233		2-Indane	Me	H	NH2	1Me-5-Ind	C		440 (M <sup>+</sup> +1)
N-g-235	ND1	N-g-191		2-Indane	Me	Me	NH2	1Me-5-1HIdz	C		455 (M <sup>+</sup> +1)
N-g-236	NA	N-g-235		2-Indane	Me	H	NH2	1Me-5-1HIdz	C		441 (M <sup>+</sup> +1)
N-g-237	ND1	N-g-193		cHex	Me	Me	NH2	2-Nap	C		417 (M <sup>+</sup> +1)
N-g-238	NA	N-g-237		cHex	Me	H	NH2	2-Nap	C		403 (M <sup>+</sup> +1)
N-g-239	ND1	N-g-195		cHex	Me	Me	NH2	1Me-5-Ind	C		420 (M <sup>+</sup> +1)
N-g-240	NA	N-g-239		cHex	Me	H	NH2	1Me-5-Ind	C		406 (M <sup>+</sup> +1)
N-g-241	ND1	N-g-197		cHex	Me	Me	NH2	5-1HIdz	C		407 (M <sup>+</sup> +1)
N-g-242	NA	N-g-241		cHex	Me	H	NH2	5-1HIdz	C		393 (M <sup>+</sup> +1)
N-g-243	ND1	N-g-199		cHex	Me	Me	NH2	1Me-5-1HIdz	C		421 (M <sup>+</sup> +1)
N-g-244	NA	N-g-243		cHex	Me	H	NH2	1Me-5-1HIdz	C		407 (M <sup>+</sup> +1)
N-g-245	ND1	N-g-201		4(Me)cHex	Me	Me	NH2	2-Nap	C		431 (M <sup>+</sup> +1)
N-g-246	NA	N-g-245		4(Me)cHex	Me	H	NH2	2-Nap	C		417 (M <sup>+</sup> +1)
N-g-247	ND1	N-g-203		4(Me)cHex	Me	Me	NH2	1Me-5-Ind	C		434 (M <sup>+</sup> +1)
N-g-248	NA	N-g-247		4(Me)cHex	Me	H	NH2	1Me-5-Ind	C		420 (M <sup>+</sup> +1)
N-g-249	ND1	N-g-205		4(Me)cHex	Me	Me	NH2	1Me-5-1HIdz	C		435 (M <sup>+</sup> +1)
N-g-250	NA	N-g-249		4(Me)cHex	Me	H	NH2	1Me-5-1HIdz	C		421 (M <sup>+</sup> +1)
N-g-251	NN1	N-g-131	CH <sub>3</sub> I	cPen	Me	Me	NHMe	2-Nap	C		417 (M <sup>+</sup> +1)
N-g-252	NA	N-g-251		cPen	Me	H	NHMe	2-Nap	C		403 (M <sup>+</sup> +1)
N-g-253	NN1	N-g-133	CH <sub>3</sub> I	cPen	Me	Me	NHMe	1Me-5-Ind	C		420 (M <sup>+</sup> +1)
N-g-254	NA	N-g-253		cPen	Me	H	NHMe	1Me-5-Ind	C		406 (M <sup>+</sup> +1)
N-g-255	NN1	N-g-137	CH <sub>3</sub> I	cPen	Me	Me	NHMe	1Me-5-1HIdz	C		421 (M <sup>+</sup> +1)
N-g-256	NA	N-g-255		cPen	Me	H	NHMe	1Me-5-1HIdz	C		407 (M <sup>+</sup> +1)
N-g-257	NN1	N-g-139	CH <sub>3</sub> I	nPr	Me	Me	NHMe	2-Nap	C		391 (M <sup>+</sup> +1)
N-g-258	NA	N-g-257		nPr	Me	H	NHMe	2-Nap	C		377 (M <sup>+</sup> +1)
N-g-259	NN1	N-g-143	CH <sub>3</sub> I	nPr	Me	Me	NHMe	1Me-5-Ind	C		394 (M <sup>+</sup> +1)
N-g-260	NA	N-g-259		nPr	Me	H	NHMe	1Me-5-Ind	C		380 (M <sup>+</sup> +1)
N-g-261	NN1	N-g-147	CH <sub>3</sub> I	nPr	Me	Me	NHMe	1Me-5-1HIdz	C		395 (M <sup>+</sup> +1)
N-g-262	NA	N-g-261		nPr	Me	H	NHMe	1Me-5-1HIdz	C		381 (M <sup>+</sup> +1)
N-g-263	NN1	N-g-149	CH <sub>3</sub> I	iPr	Me	Me	NHMe	2-Nap	C		391 (M <sup>+</sup> +1)
N-g-264	NA	N-g-263		iPr	Me	H	NHMe	2-Nap	C		377 (M <sup>+</sup> +1)
N-g-265	NN1	N-g-151	CH <sub>3</sub> I	iPr	Me	Me	NHMe	1Me-5-Ind	C		394 (M <sup>+</sup> +1)
N-g-266	NA	N-g-265		iPr	Me	H	NHMe	1Me-5-Ind	C		380 (M <sup>+</sup> +1)
N-g-267	NN1	N-g-155	CH <sub>3</sub> I	iPr	Me	Me	NHMe	1Me-5-1HIdz	C		395 (M <sup>+</sup> +1)
N-g-268	NA	N-g-267		iPr	Me	H	NHMe	1Me-5-1HIdz	C		381 (M <sup>+</sup> +1)
N-g-269	NN1	N-g-231	CH <sub>3</sub> I	2-Indane	Me	Me	NHMe	2-Nap	C		465 (M <sup>+</sup> +1)
N-g-270	NA	N-g-269		2-Indane	Me	H	NHMe	2-Nap	C		451 (M <sup>+</sup> +1)
N-g-271	NN1	N-g-233	CH <sub>3</sub> I	2-Indane	Me	Me	NHMe	1Me-5-Ind	C		468 (M <sup>+</sup> +1)
N-g-272	NA	N-g-271		2-Indane	Me	H	NHMe	1Me-5-Ind	C		454 (M <sup>+</sup> +1)
N-g-273	NN1	N-g-235	CH <sub>3</sub> I	2-Indane	Me	Me	NHMe	1Me-5-1HIdz	C		469 (M <sup>+</sup> +1)
N-g-274	NA	N-g-273		2-Indane	Me	H	NHMe	1Me-5-1HIdz	C		455 (M <sup>+</sup> +1)
N-g-275	NN1	N-g-237	CH <sub>3</sub> I	cHex	Me	Me	NHMe	2-Nap	C		431 (M <sup>+</sup> +1)
N-g-276	NA	N-g-275		cHex	Me	H	NHMe	2-Nap	C		417 (M <sup>+</sup> +1)

Tablg-N-G-7

Exp.	Syn	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									method	RTime	Mass
N-g-277	NN1	N-g-239	CH <sub>3</sub> I	cHex	Me	Me	NHMe	1Me-5-Ind	C		434 (M <sup>+</sup> +1)
N-g-278	NA	N-g-277		cHex	Me	H	NHMe	1Me-5-Ind	C		420 (M <sup>+</sup> +1)
N-g-279	NN1	N-g-245	CH <sub>3</sub> I	4Me-cHex	Me	Me	NHMe	2-Nap	C		445 (M <sup>+</sup> +1)
N-g-280	NA	N-g-279		4Me-cHex	Me	H	NHMe	2-Nap	C		431 (M <sup>+</sup> +1)
N-g-281	NN2	N-g-247	CH <sub>3</sub> I	4Me-cHex	Me	Me	NMe <sub>2</sub>	1Me-5-Ind	C		448 (M <sup>+</sup> +1)
N-g-282	NA	N-g-281		4Me-cHex	Me	H	NMe <sub>2</sub>	1Me-5-Ind	C		434 (M <sup>+</sup> +1)
N-g-283	NN2	N-g-249	CH <sub>3</sub> I	4Me-cHex	Me	Me	NMe <sub>2</sub>	1Me-5-1HIdz	C		449 (M <sup>+</sup> +1)
N-g-284	NA	N-g-283		4Me-cHex	Me	H	NMe <sub>2</sub>	1Me-5-1HIdz	C		435 (M <sup>+</sup> +1)
N-g-285	NN2	N-g-131	CH <sub>3</sub> I	cPen	Me	Me	NMe <sub>2</sub>	2-Nap	C		431 (M <sup>+</sup> +1)
N-g-286	NA	N-g-285		cPen	Me	H	NMe <sub>2</sub>	2-Nap	C		417 (M <sup>+</sup> +1)
N-g-287	NN2	N-g-133	CH <sub>3</sub> I	cPen	Me	Me	NMe <sub>2</sub>	1Me-5-Ind	C		434 (M <sup>+</sup> +1)
N-g-288	NA	N-g-287		cPen	Me	H	NMe <sub>2</sub>	1Me-5-Ind	C		420 (M <sup>+</sup> +1)
N-g-289	NN2	N-g-137	CH <sub>3</sub> I	cPen	Me	Me	NMe <sub>2</sub>	1Me-5-1HIdz	C		435 (M <sup>+</sup> +1)
N-g-290	NA	N-g-289		cPen	Me	H	NMe <sub>2</sub>	1Me-5-1HIdz	C		421 (M <sup>+</sup> +1)
N-g-291	NN2	N-g-139	CH <sub>3</sub> I	nPr	Me	Me	NMe <sub>2</sub>	2-Nap	C		405 (M <sup>+</sup> +1)
N-g-292	NA	N-g-291		nPr	Me	H	NMe <sub>2</sub>	2-Nap	C		391 (M <sup>+</sup> +1)
N-g-293	NN2	N-g-143	CH <sub>3</sub> I	nPr	Me	Me	NMe <sub>2</sub>	1Me-5-Ind	C		408 (M <sup>+</sup> +1)
N-g-294	NA	N-g-293		nPr	Me	H	NMe <sub>2</sub>	1Me-5-Ind	C		394 (M <sup>+</sup> +1)
N-g-295	NN2	N-g-147	CH <sub>3</sub> I	nPr	Me	Me	NMe <sub>2</sub>	1Me-5-1HIdz	C		409 (M <sup>+</sup> +1)
N-g-296	NA	N-g-295		nPr	Me	H	NMe <sub>2</sub>	1Me-5-1HIdz	C		395 (M <sup>+</sup> +1)
N-g-297	NN2	N-g-149	CH <sub>3</sub> I	iPr	Me	Me	NMe <sub>2</sub>	2-Nap	C		405 (M <sup>+</sup> +1)
N-g-298	NA	N-g-297		iPr	Me	H	NMe <sub>2</sub>	2-Nap	C		391 (M <sup>+</sup> +1)
N-g-299	NN2	N-g-151	CH <sub>3</sub> I	iPr	Me	Me	NMe <sub>2</sub>	1Me-5-Ind	C		408 (M <sup>+</sup> +1)
N-g-300	NA	N-g-299		iPr	Me	H	NMe <sub>2</sub>	1Me-5-Ind	C		394 (M <sup>+</sup> +1)
N-g-301	NN2	N-g-155	CH <sub>3</sub> I	iPr	Me	Me	NMe <sub>2</sub>	1Me-5-1HIdz	C		409 (M <sup>+</sup> +1)
N-g-302	NA	N-g-301		iPr	Me	H	NMe <sub>2</sub>	1Me-5-1HIdz	C		395 (M <sup>+</sup> +1)
N-g-303	NN2	N-g-231	CH <sub>3</sub> I	2Indane	Me	Me	NMe <sub>2</sub>	2-Nap	C		479 (M <sup>+</sup> +1)
N-g-304	NA	N-g-303		2Indane	Me	H	NMe <sub>2</sub>	2-Nap	C		465 (M <sup>+</sup> +1)
N-g-305	NN2	N-g-233	CH <sub>3</sub> I	2Indane	Me	Me	NMe <sub>2</sub>	1Me-5-Ind	C		482 (M <sup>+</sup> +1)
N-g-306	NA	N-g-305		2Indane	Me	H	NMe <sub>2</sub>	1Me-5-Ind	C		468 (M <sup>+</sup> +1)
N-g-307	NN2	N-g-235	CH <sub>3</sub> I	2Indane	Me	Me	NMe <sub>2</sub>	1Me-5-1HIdz	C		483 (M <sup>+</sup> +1)
N-g-308	NA	N-g-307		2Indane	Me	H	NMe <sub>2</sub>	1Me-5-1HIdz	C		469 (M <sup>+</sup> +1)
N-g-309	NN2	N-g-237	CH <sub>3</sub> I	cHex	Me	Me	NMe <sub>2</sub>	2-Nap	C		445 (M <sup>+</sup> +1)
N-g-310	NA	N-g-265		cHex	Me	H	NMe <sub>2</sub>	2-Nap	C		431 (M <sup>+</sup> +1)
N-g-311	NN2	N-g-239	CH <sub>3</sub> I	cHex	Me	Me	NMe <sub>2</sub>	1Me-5-Ind	C		448 (M <sup>+</sup> +1)
N-g-312	NA	N-g-267		cHex	Me	H	NMe <sub>2</sub>	1Me-5-Ind	C		434 (M <sup>+</sup> +1)
N-g-313	NN2	N-g-245	CH <sub>3</sub> I	4Me-cHex	Me	Me	NMe <sub>2</sub>	2-Nap	C		459 (M <sup>+</sup> +1)
N-g-314	NA	N-g-269		4Me-cHex	Me	H	NMe <sub>2</sub>	2-Nap	C		445 (M <sup>+</sup> +1)
N-g-315	NN2	N-g-247	CH <sub>3</sub> I	4Me-cHex	Me	Me	NMe <sub>2</sub>	1Me-5-Ind	C		462 (M <sup>+</sup> +1)
N-g-316	NA	N-g-271		4Me-cHex	Me	H	NMe <sub>2</sub>	1Me-5-Ind	C		448 (M <sup>+</sup> +1)
N-g-317	NN2	N-g-249	CH <sub>3</sub> I	4Me-cHex	Me	Me	NMe <sub>2</sub>	1Me-5-1HIdz	C		463 (M <sup>+</sup> +1)
N-g-318	NA	N-g-273		4Me-cHex	Me	H	NMe <sub>2</sub>	1Me-5-1HIdz	C		449 (M <sup>+</sup> +1)

[Examples N-h-1 to N-h-458]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification are shown in Table-N-H-1 to Table-N-H-10. In the tables, the compound numbers are mentioned in the columns indicated as "Exp.". In the tables, corresponding methods among the aforementioned synthesis methods are shown in the columns of "Syn" with symbols,

the starting compounds 1 are mentioned in the columns of "SM1", and the starting compounds 2 are mentioned in the columns of "SM2".

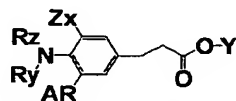


Table-N-H-1

Exp.	Syn	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									method	RTime	Mass
N-h-1	NB1	Int.n-89	BRA1	Bn	H	Me	NO2	2-Nap	C		441 (M <sup>+</sup> +1)
N-h-2	NA	N-h-1		Bn	H	H	NO2	2-Nap	C		427 (M <sup>+</sup> +1)
N-h-3	NB1	Int.n-89	BRA2	Bn	H	Me	NO2	5-Ind	C		430 (M <sup>+</sup> +1)
N-h-4	NA	N-h-3		Bn	H	H	NO2	5-Ind	C		416 (M <sup>+</sup> +1)
N-h-5	NB1	Int.n-89	BRA3	Bn	H	Me	NO2	1Me-5-Ind	C		444 (M <sup>+</sup> +1)
N-h-6	NA	N-h-5		Bn	H	H	NO2	1Me-5-Ind	C		430 (M <sup>+</sup> +1)
N-h-7	NB1	Int.n-89	BRA5	Bn	H	Me	NO2	5-1HIdz	C		431 (M <sup>+</sup> +1)
N-h-8	NA	N-h-7		Bn	H	H	NO2	5-1HIdz	C		417 (M <sup>+</sup> +1)
N-h-9	NB1	Int.n-89	BRA6	Bn	H	Me	NO2	1Me-5-1HIdz	C		445 (M <sup>+</sup> +1)
N-h-10	NA	N-h-9		Bn	H	H	NO2	1Me-5-1HIdz	C		431 (M <sup>+</sup> +1)
N-h-11	NB1	Int.n-89	BRA10	Bn	H	Me	NO2	3-Qu	C		442 (M <sup>+</sup> +1)
N-h-12	NA	N-h-11		Bn	H	H	NO2	3-Qu	C		428 (M <sup>+</sup> +1)
N-h-13	NB1	Int.n-89	BRA11	Bn	H	Me	NO2	6-Qu	C		442 (M <sup>+</sup> +1)
N-h-14	NA	N-h-13		Bn	H	H	NO2	6-Qu	C		428 (M <sup>+</sup> +1)
N-h-15	NB1	Int.n-89	BRA12	Bn	H	Me	NO2	6-IQ	C		442 (M <sup>+</sup> +1)
N-h-16	NA	N-h-15		Bn	H	H	NO2	6-IQ	C		428 (M <sup>+</sup> +1)
N-h-17	NB1	Int.n-90	BRA1	4FBn	H	Me	NO2	2-Nap	C		459 (M <sup>+</sup> +1)
N-h-18	NA	N-h-17		4FBn	H	H	NO2	2-Nap	C		445 (M <sup>+</sup> +1)
N-h-19	NB1	Int.n-90	BRA2	4FBn	H	Me	NO2	5-Ind	C		448 (M <sup>+</sup> +1)
N-h-20	NA	N-h-19		4FBn	H	H	NO2	5-Ind	C		434 (M <sup>+</sup> +1)
N-h-21	NB1	Int.n-90	BRA3	4FBn	H	Me	NO2	1Me-5-Ind	C		462 (M <sup>+</sup> +1)
N-h-22	NA	N-h-21		4FBn	H	H	NO2	1Me-5-Ind	C		448 (M <sup>+</sup> +1)
N-h-23	NB1	Int.n-90	BRA5	4FBn	H	Me	NO2	5-1HIdz	C		449 (M <sup>+</sup> +1)
N-h-24	NA	N-h-23		4FBn	H	H	NO2	5-1HIdz	C		435 (M <sup>+</sup> +1)
N-h-25	NB1	Int.n-90	BRA6	4FBn	H	Me	NO2	1Me-5-1HIdz	C		463 (M <sup>+</sup> +1)
N-h-26	NA	N-h-25		4FBn	H	H	NO2	1Me-5-1HIdz	C		449 (M <sup>+</sup> +1)
N-h-27	NB1	Int.n-91	BRA2	2FBn	H	Me	NO2	5-Ind	C		448 (M <sup>+</sup> +1)
N-h-28	NA	N-h-27		2FBn	H	H	NO2	5-Ind	C		434 (M <sup>+</sup> +1)
N-h-29	NB1	Int.n-91	BRA3	2FBn	H	Me	NO2	1Me-5-Ind	C		462 (M <sup>+</sup> +1)
N-h-30	NA	N-h-29		2FBn	H	H	NO2	1Me-5-Ind	C		448 (M <sup>+</sup> +1)
N-h-31	NB1	Int.n-91	BRA5	2FBn	H	Me	NO2	5-1HIdz	C		449 (M <sup>+</sup> +1)
N-h-32	NA	N-h-31		2FBn	H	H	NO2	5-1HIdz	C		435 (M <sup>+</sup> +1)
N-h-33	NB1	Int.n-91	BRA6	2FBn	H	Me	NO2	1Me-5-1HIdz	C		463 (M <sup>+</sup> +1)
N-h-34	NA	N-h-33		2FBn	H	H	NO2	1Me-5-1HIdz	C		449 (M <sup>+</sup> +1)
N-h-35	NB1	Int.n-92	BRA2	3FBn	H	Me	NO2	5-Ind	C		448 (M <sup>+</sup> +1)
N-h-36	NA	N-h-35		3FBn	H	H	NO2	5-Ind	C		434 (M <sup>+</sup> +1)
N-h-37	NB1	Int.n-92	BRA3	3FBn	H	Me	NO2	1Me-5-Ind	C		462 (M <sup>+</sup> +1)
N-h-38	NA	N-h-37		3FBn	H	H	NO2	1Me-5-Ind	C		448 (M <sup>+</sup> +1)
N-h-39	NB1	Int.n-92	BRA5	3FBn	H	Me	NO2	5-1HIdz	C		449 (M <sup>+</sup> +1)
N-h-40	NA	N-h-39		3FBn	H	H	NO2	5-1HIdz	C		435 (M <sup>+</sup> +1)
N-h-41	NB1	Int.n-92	BRA6	3FBn	H	Me	NO2	1Me-5-1HIdz	C		463 (M <sup>+</sup> +1)
N-h-42	NA	N-h-41		3FBn	H	H	NO2	1Me-5-1HIdz	C		449 (M <sup>+</sup> +1)
N-h-43	NB1	Int.n-93	BRA1	2,3DFBn	H	Me	NO2	2-Nap	C		477 (M <sup>+</sup> +1)
N-h-44	NA	N-h-43		2,3DFBn	H	H	NO2	2-Nap	C		463 (M <sup>+</sup> +1)
N-h-45	NB1	Int.n-93	BRA3	2,3DFBn	H	Me	NO2	1Me-5-Ind	C		480 (M <sup>+</sup> +1)
N-h-46	NA	N-h-45		2,3DFBn	H	H	NO2	1Me-5-Ind	C		466 (M <sup>+</sup> +1)

Table-N-H-2

Exp.	Syn	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									method	RTime	Mass
N-h-47	NB1	Intn-93	BRA5	2,3DFBn	H	Me	NO2	5-1HIdz	C		467 (M <sup>+</sup> +1)
N-h-48	NA	N-h-47		2,3DFBn	H	H	NO2	5-1HIdz	C		453 (M <sup>+</sup> +1)
N-h-49	NB1	Intn-93	BRA6	2,3DFBn	H	Me	NO2	1Me-5-1HIdz	C		481 (M <sup>+</sup> +1)
N-h-50	NA	N-h-49		2,3DFBn	H	H	NO2	1Me-5-1HIdz	C		467 (M <sup>+</sup> +1)
N-h-51	NB1	Intn-94	BRA2	3,4DFBn	H	Me	NO2	5-Ind	C		466 (M <sup>+</sup> +1)
N-h-52	NA	N-h-51		3,4DFBn	H	H	NO2	5-Ind	C		452 (M <sup>+</sup> +1)
N-h-53	NB1	Intn-94	BRA3	3,4DFBn	H	Me	NO2	1Me-5-Ind	C		480 (M <sup>+</sup> +1)
N-h-54	NA	N-h-53		3,4DFBn	H	H	NO2	1Me-5-Ind	C		466 (M <sup>+</sup> +1)
N-h-55	NB1	Intn-94	BRA6	3,4DFBn	H	Me	NO2	1Me-5-1HIdz	C		481 (M <sup>+</sup> +1)
N-h-56	NA	N-h-55		3,4DFBn	H	H	NO2	1Me-5-1HIdz	C		467 (M <sup>+</sup> +1)
N-h-57	NB1	Intn-95	BRA1	4PhBn	H	Me	NO2	2-Nap	C		517 (M <sup>+</sup> +1)
N-h-58	NA	N-h-57		4PhBn	H	H	NO2	2-Nap	C		503 (M <sup>+</sup> +1)
N-h-59	NB1	Intn-95	BRA2	4PhBn	H	Me	NO2	5-Ind	C		506 (M <sup>+</sup> +1)
N-h-60	NA	N-h-59		4PhBn	H	H	NO2	5-Ind	C		492 (M <sup>+</sup> +1)
N-h-61	NB1	Intn-95	BRA3	4PhBn	H	Me	NO2	1Me-5-Ind	C		520 (M <sup>+</sup> +1)
N-h-62	NA	N-h-61		4PhBn	H	H	NO2	1Me-5-Ind	C		506 (M <sup>+</sup> +1)
N-h-63	NB1	Intn-95	BRA5	4PhBn	H	Me	NO2	5-1HIdz	C		507 (M <sup>+</sup> +1)
N-h-64	NA	N-h-63		4PhBn	H	H	NO2	5-1HIdz	C		493 (M <sup>+</sup> +1)
N-h-65	NB1	Intn-96	BRA1	2CF3Bn	H	Me	NO2	2-Nap	C		509 (M <sup>+</sup> +1)
N-h-66	NA	N-h-65		2CF3Bn	H	H	NO2	2-Nap	C		495 (M <sup>+</sup> +1)
N-h-67	NB1	Intn-96	BRA2	2CF3Bn	H	Me	NO2	5-Ind	C		498 (M <sup>+</sup> +1)
N-h-68	NA	N-h-67		2CF3Bn	H	H	NO2	5-Ind	C		484 (M <sup>+</sup> +1)
N-h-69	NB1	Intn-96	BRA3	2CF3Bn	H	Me	NO2	1Me-5-Ind	C		512 (M <sup>+</sup> +1)
N-h-70	NA	N-h-69		2CF3Bn	H	H	NO2	1Me-5-Ind	C		498 (M <sup>+</sup> +1)
N-h-71	NB1	Intn-96	BRA5	2CF3Bn	H	Me	NO2	5-1HIdz	C		499 (M <sup>+</sup> +1)
N-h-72	NA	N-h-71		2CF3Bn	H	H	NO2	5-1HIdz	C		485 (M <sup>+</sup> +1)
N-h-73	NB1	Intn-97	BRA2	2-TF	H	Me	NO2	5-Ind	C		436 (M <sup>+</sup> +1)
N-h-74	NA	N-h-73		2-TF	H	H	NO2	5-Ind	C		422 (M <sup>+</sup> +1)
N-h-75	NB1	Intn-97	BRA3	2-TF	H	Me	NO2	1Me-5-Ind	C		450 (M <sup>+</sup> +1)
N-h-76	NA	N-h-75		2-TF	H	H	NO2	1Me-5-Ind	C		436 (M <sup>+</sup> +1)
N-h-77	NB1	Intn-97	BRA6	2-TF	H	Me	NO2	1Me-5-1HIdz	C		451 (M <sup>+</sup> +1)
N-h-78	NA	N-h-77		2-TF	H	H	NO2	1Me-5-1HIdz	C		437 (M <sup>+</sup> +1)
N-h-79	NB1	Intn-98	BRA2	3-TF	H	Me	NO2	5-Ind	C		436 (M <sup>+</sup> +1)
N-h-80	NA	N-h-79		3-TF	H	H	NO2	5-Ind	C		422 (M <sup>+</sup> +1)
N-h-81	NB1	Intn-98	BRA3	3-TF	H	Me	NO2	1Me-5-Ind	C		450 (M <sup>+</sup> +1)
N-h-82	NA	N-h-81		3-TF	H	H	NO2	1Me-5-Ind	C		436 (M <sup>+</sup> +1)
N-h-83	NB1	Intn-98	BRA5	3-TF	H	Me	NO2	5-1HIdz	C		437 (M <sup>+</sup> +1)
N-h-84	NA	N-h-83		3-TF	H	H	NO2	5-1HIdz	C		423 (M <sup>+</sup> +1)
N-h-85	NB1	Intn-98	BRA6	3-TF	H	Me	NO2	1Me-5-1HIdz	C		451 (M <sup>+</sup> +1)
N-h-86	NA	N-h-85		3-TF	H	H	NO2	1Me-5-1HIdz	C		437 (M <sup>+</sup> +1)
N-h-87	NB1	Intn-99	BRA1	2-FR	H	Me	NO2	2Nap	C		459 (M <sup>+</sup> +1)
N-h-88	NA	N-h-87		2-FR	H	H	NO2	2Nap	C		445 (M <sup>+</sup> +1)
N-h-89	NB1	Intn-99	BRA2	2-FR	H	Me	NO2	5-Ind	C		420 (M <sup>+</sup> +1)
N-h-90	NA	N-h-89		2-FR	H	H	NO2	5-Ind	C		406 (M <sup>+</sup> +1)
N-h-91	NB1	Intn-99	BRA6	2-FR	H	Me	NO2	1Me-5-1HIdz	C		434 (M <sup>+</sup> +1)
N-h-92	NA	N-h-91		2-FR	H	H	NO2	1Me-5-1HIdz	C		420 (M <sup>+</sup> +1)

Table-N-H-3

Exp.	Syn	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									method	RTime	Mass
N-h-93	NB1	Intn-100	BRA1	Bn	Me	Me	NO2	2-Nap	C		455 (M <sup>+</sup> +1)
N-h-94	NA	N-h-93		Bn	Me	H	NO2	2-Nap	C		427 (M <sup>+</sup> +1)
N-h-95	NB1	Intn-100	BRA2	Bn	Me	Me	NO2	5-Ind	C		430 (M <sup>+</sup> +1)
N-h-96	NA	N-h-95		Bn	Me	H	NO2	5-Ind	C		416 (M <sup>+</sup> +1)
N-h-97	NB1	Intn-100	BRA3	Bn	Me	Me	NO2	1Me-5-Ind	C		444 (M <sup>+</sup> +1)
N-h-98	NA	N-h-97		Bn	Me	H	NO2	1Me-5-Ind	C		430 (M <sup>+</sup> +1)
N-h-99	NB1	Intn-100	BRA5	Bn	Me	Me	NO2	5-1HIdz	C		431 (M <sup>+</sup> +1)
N-h-100	NA	N-h-99		Bn	Me	H	NO2	5-1HIdz	C		417 (M <sup>+</sup> +1)
N-h-101	NB1	Intn-100	BRA6	Bn	Me	Me	NO2	1Me-5-1HIdz	C		445 (M <sup>+</sup> +1)
N-h-102	NA	N-h-101		Bn	Me	H	NO2	1Me-5-1HIdz	C		431 (M <sup>+</sup> +1)
N-h-103	NB1	Intn-100	BRA10	Bn	Me	Me	NO2	3-Qu	C		442 (M <sup>+</sup> +1)
N-h-104	NA	N-h-103		Bn	Me	H	NO2	3-Qu	C		428 (M <sup>+</sup> +1)
N-h-105	NB1	Intn-100	BRA11	Bn	Me	Me	NO2	6-Qu	C		442 (M <sup>+</sup> +1)
N-h-106	NA	N-h-105		Bn	Me	H	NO2	6-Qu	C		428 (M <sup>+</sup> +1)
N-h-107	NB1	Intn-100	BRA12	Bn	Me	Me	NO2	6-IQ	C		442 (M <sup>+</sup> +1)
N-h-108	NA	N-h-107		Bn	Me	H	NO2	6-IQ	C		428 (M <sup>+</sup> +1)
N-h-109	NB1	Intn-101	BRA1	4FBn	Me	Me	NO2	2-Nap	C		459 (M <sup>+</sup> +1)
N-h-110	NA	N-h-109		4FBn	Me	H	NO2	2-Nap	C		445 (M <sup>+</sup> +1)
N-h-111	NB1	Intn-101	BRA2	4FBn	Me	Me	NO2	5-Ind	C		448 (M <sup>+</sup> +1)
N-h-112	NA	N-h-111		4FBn	Me	H	NO2	5-Ind	C		434 (M <sup>+</sup> +1)
N-h-113	NB1	Intn-101	BRA3	4FBn	Me	Me	NO2	1Me-5-Ind	C		462 (M <sup>+</sup> +1)
N-h-114	NA	N-h-113		4FBn	Me	H	NO2	1Me-5-Ind	C		448 (M <sup>+</sup> +1)
N-h-115	NB1	Intn-101	BRA5	4FBn	Me	Me	NO2	5-1HIdz	C		449 (M <sup>+</sup> +1)
N-h-116	NA	N-h-115		4FBn	Me	H	NO2	5-1HIdz	C		435 (M <sup>+</sup> +1)
N-h-117	NB1	Intn-101	BRA6	4FBn	Me	Me	NO2	1Me-5-1HIdz	C		463 (M <sup>+</sup> +1)
N-h-118	NA	N-h-117		4FBn	Me	H	NO2	1Me-5-1HIdz	C		449 (M <sup>+</sup> +1)
N-h-119	NB1	Intn-102	BRA1	2FBn	Me	Me	NO2	2-Nap	C		448 (M <sup>+</sup> +1)
N-h-120	NA	N-h-119		2FBn	Me	H	NO2	2-Nap	C		434 (M <sup>+</sup> +1)
N-h-121	NB1	Intn-102	BRA3	2FBn	Me	Me	NO2	1Me-5-Ind	C		462 (M <sup>+</sup> +1)
N-h-122	NA	N-h-121		2FBn	Me	H	NO2	1Me-5-Ind	C		448 (M <sup>+</sup> +1)
N-h-123	NB1	Intn-102	BRA5	2FBn	Me	Me	NO2	5-1HIdz	C		449 (M <sup>+</sup> +1)
N-h-124	NA	N-h-123		2FBn	Me	H	NO2	5-1HIdz	C		435 (M <sup>+</sup> +1)
N-h-125	NB1	Intn-102	BRA6	2FBn	Me	Me	NO2	1Me-5-1HIdz	C		463 (M <sup>+</sup> +1)
N-h-126	NA	N-h-125		2FBn	Me	H	NO2	1Me-5-1HIdz	C		449 (M <sup>+</sup> +1)
N-h-127	NB1	Intn-103	BRA1	3FBn	Me	Me	NO2	2-Nap	C		448 (M <sup>+</sup> +1)
N-h-128	NA	N-h-127		3FBn	Me	H	NO2	2-Nap	C		434 (M <sup>+</sup> +1)
N-h-129	NB1	Intn-103	BRA3	3FBn	Me	Me	NO2	1Me-5-Ind	C		462 (M <sup>+</sup> +1)
N-h-130	NA	N-h-129		3FBn	Me	H	NO2	1Me-5-Ind	C		448 (M <sup>+</sup> +1)
N-h-131	NB1	Intn-103	BRA5	3FBn	Me	Me	NO2	5-1HIdz	C		449 (M <sup>+</sup> +1)
N-h-132	NA	N-h-131		3FBn	Me	H	NO2	5-1HIdz	C		435 (M <sup>+</sup> +1)
N-h-133	NB1	Intn-103	BRA6	3FBn	Me	Me	NO2	1Me-5-1HIdz	C		463 (M <sup>+</sup> +1)
N-h-134	NA	N-h-133		3FBn	Me	H	NO2	1Me-5-1HIdz	C		449 (M <sup>+</sup> +1)
N-h-135	NB1	Intn-104	BRA1	2,3DFBn	Me	Me	NO2	2-Nap	C		477 (M <sup>+</sup> +1)
N-h-136	NA	N-h-135		2,3DFBn	Me	H	NO2	2-Nap	C		463 (M <sup>+</sup> +1)
N-h-137	NB1	Intn-104	BRA3	2,3DFBn	Me	Me	NO2	1Me-5-Ind	C		480 (M <sup>+</sup> +1)
N-h-138	NA	N-h-137		2,3DFBn	Me	H	NO2	1Me-5-Ind	C		466 (M <sup>+</sup> +1)

Table-N-H-4

Exp.	Syn	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									method	RTime	Mass
N-h-139	NB1	Int.n-104	BRA5	2,3DFBn	Me	Me	NO2	5-1HIdz	C		481 (M <sup>+</sup> +1)
N-h-140	NA	N-h-139		2,3DFBn	Me	H	NO2	5-1HIdz	C		467 (M <sup>+</sup> +1)
N-h-141	NB1	Int.n-104	BRA6	2,3DFBn	Me	Me	NO2	1Me-5-1HIdz	C		495 (M <sup>+</sup> +1)
N-h-142	NA	N-h-141		2,3DFBn	Me	H	NO2	1Me-5-1HIdz	C		481 (M <sup>+</sup> +1)
N-h-143	NB1	Int.n-105	BRA1	3,4DFBn	Me	Me	NO2	2-Nap	C		480 (M <sup>+</sup> +1)
N-h-144	NA	N-h-143		3,4DFBn	Me	H	NO2	2-Nap	C		466 (M <sup>+</sup> +1)
N-h-145	NB1	Int.n-105	BRA3	3,4DFBn	Me	Me	NO2	1Me-5-Ind	C		494 (M <sup>+</sup> +1)
N-h-146	NA	N-h-145		3,4DFBn	Me	H	NO2	1Me-5-Ind	C		480 (M <sup>+</sup> +1)
N-h-147	NB1	Int.n-105	BRA6	3,4DFBn	Me	Me	NO2	1Me-5-1HIdz	C		495 (M <sup>+</sup> +1)
N-h-148	NA	N-h-147		3,4DFBn	Me	H	NO2	1Me-5-1HIdz	C		481 (M <sup>+</sup> +1)
N-h-149	NB1	Int.n-106	BRA1	4PhBn	Me	Me	NO2	2-Nap	C		531 (M <sup>+</sup> +1)
N-h-150	NA	N-h-149		4PhBn	Me	H	NO2	2-Nap	C		517 (M <sup>+</sup> +1)
N-h-151	NB1	Int.n-106	BRA2	4PhBn	Me	Me	NO2	5-Ind	C		520 (M <sup>+</sup> +1)
N-h-152	NA	N-h-151		4PhBn	Me	H	NO2	5-Ind	C		506 (M <sup>+</sup> +1)
N-h-153	NB1	Int.n-106	BRA3	4PhBn	Me	Me	NO2	1Me-5-Ind	C		534 (M <sup>+</sup> +1)
N-h-154	NA	N-h-153		4PhBn	Me	H	NO2	1Me-5-Ind	C		520 (M <sup>+</sup> +1)
N-h-155	NB1	Int.n-106	BRA6	4PhBn	Me	Me	NO2	1Me-5-1HIdz	C		521 (M <sup>+</sup> +1)
N-h-156	NA	N-h-155		4PhBn	Me	H	NO2	1Me-5-1HIdz	C		507 (M <sup>+</sup> +1)
N-h-157	NB1	Int.n-107	BRA1	2CF3Bn	Me	Me	NO2	2-Nap	C		523 (M <sup>+</sup> +1)
N-h-158	NA	N-h-157		2CF3Bn	Me	H	NO2	2-Nap	C		509 (M <sup>+</sup> +1)
N-h-159	NB1	Int.n-107	BRA2	2CF3Bn	Me	Me	NO2	5-Ind	C		512 (M <sup>+</sup> +1)
N-h-160	NA	N-h-159		2CF3Bn	Me	H	NO2	5-Ind	C		498 (M <sup>+</sup> +1)
N-h-161	NB1	Int.n-107	BRA3	2CF3Bn	Me	Me	NO2	1Me-5-Ind	C		526 (M <sup>+</sup> +1)
N-h-162	NA	N-h-161		2CF3Bn	Me	H	NO2	1Me-5-Ind	C		512 (M <sup>+</sup> +1)
N-h-163	NB1	Int.n-107	BRA5	2CF3Bn	Me	Me	NO2	5-1HIdz	C		513 (M <sup>+</sup> +1)
N-h-164	NA	N-h-163		2CF3Bn	Me	H	NO2	5-1HIdz	C		499 (M <sup>+</sup> +1)
N-h-165	NB1	Int.n-108	BRA1	2-TF	Me	Me	NO2	2-Nap	C		450 (M <sup>+</sup> +1)
N-h-166	NA	N-h-165		2-TF	Me	H	NO2	2-Nap	C		436 (M <sup>+</sup> +1)
N-h-167	NB1	Int.n-108	BRA3	2-TF	Me	Me	NO2	1Me-5-Ind	C		464 (M <sup>+</sup> +1)
N-h-168	NA	N-h-167		2-TF	Me	H	NO2	1Me-5-Ind	C		450 (M <sup>+</sup> +1)
N-h-169	NB1	Int.n-108	BRA6	2-TF	Me	Me	NO2	1Me-5-1HIdz	C		465 (M <sup>+</sup> +1)
N-h-170	NA	N-h-169		2-TF	Me	H	NO2	1Me-5-1HIdz	C		451 (M <sup>+</sup> +1)
N-h-171	NB1	Int.n-109	BRA1	3-TF	Me	Me	NO2	2-Nap	C		450 (M <sup>+</sup> +1)
N-h-172	NA	N-h-171		3-TF	Me	H	NO2	2-Nap	C		436 (M <sup>+</sup> +1)
N-h-173	NB1	Int.n-109	BRA2	3-TF	Me	Me	NO2	5-Ind	C		464 (M <sup>+</sup> +1)
N-h-174	NA	N-h-173		3-TF	Me	H	NO2	5-Ind	C		450 (M <sup>+</sup> +1)
N-h-175	NB1	Int.n-109	BRA3	3-TF	Me	Me	NO2	1Me-5-Ind	C		451 (M <sup>+</sup> +1)
N-h-176	NA	N-h-175		3-TF	Me	H	NO2	1Me-5-Ind	C		437 (M <sup>+</sup> +1)
N-h-177	NB1	Int.n-110	BRA6	3-TF	Me	Me	NO2	1Me-5-1HIdz	C		465 (M <sup>+</sup> +1)
N-h-178	NA	N-h-177		3-TF	Me	H	NO2	1Me-5-1HIdz	C		451 (M <sup>+</sup> +1)
N-h-179	NB1	Int.n-110	BRA1	2-FR	Me	Me	NO2	2-Nap	C		473 (M <sup>+</sup> +1)
N-h-180	NA	N-h-179		2-FR	Me	H	NO2	2-Nap	C		459 (M <sup>+</sup> +1)
N-h-181	NB1	Int.n-110	BRA2	2-FR	Me	Me	NO2	5-Ind	C		434 (M <sup>+</sup> +1)
N-h-182	NA	N-h-181		2-FR	Me	H	NO2	5-Ind	C		420 (M <sup>+</sup> +1)
N-h-183	NB1	Int.n-109	BRA6	2-FR	Me	Me	NO2	1Me-5-1HIdz	C		448 (M <sup>+</sup> +1)
N-h-184	NA	N-h-183		2-FR	Me	H	NO2	1Me-5-1HIdz	C		434 (M <sup>+</sup> +1)

Table-N-H-5

Exp.	Syn	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									method	RTime	Mass
N-h-185	ND1	N-h-1		Bn	H	Me	NH2	2-Nap	C		411 (M <sup>+</sup> +1)
N-h-186	NA	N-h-185		Bn	H	H	NH2	2-Nap	C		397 (M <sup>+</sup> +1)
N-h-187	ND1	N-h-3		Bn	H	Me	NH2	5-Ind	C		400 (M <sup>+</sup> +1)
N-h-188	NA	N-h-187		Bn	H	H	NH2	5-Ind	C		386 (M <sup>+</sup> +1)
N-h-189	ND1	N-h-5		Bn	H	Me	NH2	1Me-5-Ind	C		414 (M <sup>+</sup> +1)
N-h-190	NA	N-h-189		Bn	H	H	NH2	1Me-5-Ind	C		400 (M <sup>+</sup> +1)
N-h-191	ND1	N-h-7		Bn	H	Me	NH2	5-1HIdz	C		401 (M <sup>+</sup> +1)
N-h-192	NA	N-h-191		Bn	H	H	NH2	5-1HIdz	C		387 (M <sup>+</sup> +1)
N-h-193	ND1	N-h-9		Bn	H	Me	NH2	1Me-5-1HIdz	C		415 (M <sup>+</sup> +1)
N-h-194	NA	N-h-193		Bn	H	H	NH2	1Me-5-1HIdz	C		401 (M <sup>+</sup> +1)
N-h-195	ND1	N-h-11		Bn	H	Me	NH2	3-Qu	C		412 (M <sup>+</sup> +1)
N-h-196	NA	N-h-195		Bn	H	H	NH2	3-Qu	C		398 (M <sup>+</sup> +1)
N-h-197	ND1	N-h-13		Bn	H	Me	NH2	6-Qu	C		412 (M <sup>+</sup> +1)
N-h-198	NA	N-h-197		Bn	H	H	NH2	6-Qu	C		398 (M <sup>+</sup> +1)
N-h-199	ND1	N-h-17		4FBn	H	Me	NH2	2-Nap	C		429 (M <sup>+</sup> +1)
N-h-200	NA	N-h-199		4FBn	H	H	NH2	2-Nap	C		415 (M <sup>+</sup> +1)
N-h-201	ND1	N-h-19		4FBn	H	Me	NH2	5-Ind	C		418 (M <sup>+</sup> +1)
N-h-202	NA	N-h-201		4FBn	H	H	NH2	5-Ind	C		404 (M <sup>+</sup> +1)
N-h-203	ND1	N-h-21		4FBn	H	Me	NH2	1Me-5-Ind	C		432 (M <sup>+</sup> +1)
N-h-204	NA	N-h-203		4FBn	H	H	NH2	1Me-5-Ind	C		418 (M <sup>+</sup> +1)
N-h-205	ND1	N-h-23		4FBn	H	Me	NH2	5-1HIdz	C		419 (M <sup>+</sup> +1)
N-h-206	NA	N-h-205		4FBn	H	H	NH2	5-1HIdz	C		405 (M <sup>+</sup> +1)
N-h-207	ND1	N-h-25		4FBn	H	Me	NH2	1Me-5-1HIdz	C		433 (M <sup>+</sup> +1)
N-h-208	NA	N-h-207		4FBn	H	H	NH2	1Me-5-1HIdz	C		419 (M <sup>+</sup> +1)
N-h-209	ND1	N-h-27		2FBn	H	Me	NH2	5-Ind	C		418 (M <sup>+</sup> +1)
N-h-210	NA	N-h-209		2FBn	H	H	NH2	5-Ind	C		404 (M <sup>+</sup> +1)
N-h-211	ND1	N-h-29		2FBn	H	Me	NH2	1Me-5-Ind	C		432 (M <sup>+</sup> +1)
N-h-212	NA	N-h-211		2FBn	H	H	NH2	1Me-5-Ind	C		418 (M <sup>+</sup> +1)
N-h-213	ND1	N-h-31		2FBn	H	Me	NH2	5-1HIdz	C		419 (M <sup>+</sup> +1)
N-h-214	NA	N-h-213		2FBn	H	H	NH2	5-1HIdz	C		405 (M <sup>+</sup> +1)
N-h-215	ND1	N-h-33		2FBn	H	Me	NH2	1Me-5-1HIdz	C		433 (M <sup>+</sup> +1)
N-h-216	NA	N-h-215		2FBn	H	H	NH2	1Me-5-1HIdz	C		419 (M <sup>+</sup> +1)
N-h-217	ND1	N-h-35		3FBn	H	Me	NH2	5-Ind	C		418 (M <sup>+</sup> +1)
N-h-218	NA	N-h-217		3FBn	H	H	NH2	5-Ind	C		404 (M <sup>+</sup> +1)
N-h-219	ND1	N-h-37		3FBn	H	Me	NH2	1Me-5-Ind	C		432 (M <sup>+</sup> +1)
N-h-220	NA	N-h-219		3FBn	H	H	NH2	1Me-5-Ind	C		418 (M <sup>+</sup> +1)
N-h-221	ND1	N-h-39		3FBn	H	Me	NH2	5-1HIdz	C		419 (M <sup>+</sup> +1)
N-h-222	NA	N-h-221		3FBn	H	H	NH2	5-1HIdz	C		405 (M <sup>+</sup> +1)
N-h-223	ND1	N-h-41		3FBn	H	Me	NH2	1Me-5-1HIdz	C		433 (M <sup>+</sup> +1)
N-h-224	NA	N-h-223		3FBn	H	H	NH2	1Me-5-1HIdz	C		419 (M <sup>+</sup> +1)
N-h-225	ND1	N-h-43		2,3DFBn	H	Me	NH2	2-Nap	C		447 (M <sup>+</sup> +1)
N-h-226	NA	N-h-225		2,3DFBn	H	H	NH2	2-Nap	C		433 (M <sup>+</sup> +1)
N-h-227	ND1	N-h-45		2,3DFBn	H	Me	NH2	1Me-5-Ind	C		450 (M <sup>+</sup> +1)
N-h-228	NA	N-h-227		2,3DFBn	H	H	NH2	1Me-5-Ind	C		436 (M <sup>+</sup> +1)
N-h-229	ND1	N-h-47		2,3DFBn	H	Me	NH2	5-1HIdz	C		437 (M <sup>+</sup> +1)
N-h-230	NA	N-h-229		2,3DFBn	H	H	NH2	5-1HIdz	C		423 (M <sup>+</sup> +1)

Table-N-H-6

Exp.	Syn	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									method	RTime	Mass
N-h-231	ND1	N-h-49		2,3DFBn	H	Me	NH2	1Me-5-1HIdz	C		451 (M <sup>+</sup> +1)
N-h-232	NA	N-h-231		2,3DFBn	H	H	NH2	1Me-5-1HIdz	C		437 (M <sup>+</sup> +1)
N-h-233	ND1	N-h-51		3,4DFBn	H	Me	NH2	5-Ind	C		436 (M <sup>+</sup> +1)
N-h-234	NA	N-h-233		3,4DFBn	H	H	NH2	5-Ind	C		422 (M <sup>+</sup> +1)
N-h-235	ND1	N-h-53		3,4DFBn	H	Me	NH2	1Me-5-Ind	C		450 (M <sup>+</sup> +1)
N-h-236	NA	N-h-235		3,4DFBn	H	H	NH2	1Me-5-Ind	C		436 (M <sup>+</sup> +1)
N-h-237	ND1	N-h-55		3,4DFBn	H	Me	NH2	1Me-5-1HIdz	C		451 (M <sup>+</sup> +1)
N-h-238	NA	N-h-237		3,4DFBn	H	H	NH2	1Me-5-1HIdz	C		437 (M <sup>+</sup> +1)
N-h-239	ND1	N-h-57		4PhBn	H	Me	NH2	2-Nap	C		487 (M <sup>+</sup> +1)
N-h-240	NA	N-h-239		4PhBn	H	H	NH2	2-Nap	C		473 (M <sup>+</sup> +1)
N-h-241	ND1	N-h-59		4PhBn	H	Me	NH2	5-Ind	C		476 (M <sup>+</sup> +1)
N-h-242	NA	N-h-241		4PhBn	H	H	NH2	5-Ind	C		462 (M <sup>+</sup> +1)
N-h-243	ND1	N-h-61		4PhBn	H	Me	NH2	1Me-5-Ind	C		490 (M <sup>+</sup> +1)
N-h-244	NA	N-h-243		4PhBn	H	H	NH2	1Me-5-Ind	C		476 (M <sup>+</sup> +1)
N-h-245	ND1	N-h-63		4PhBn	H	Me	NH2	5-1HIdz	C		477 (M <sup>+</sup> +1)
N-h-246	NA	N-h-245		4PhBn	H	H	NH2	5-1HIdz	C		463 (M <sup>+</sup> +1)
N-h-247	ND1	N-h-65		2CF3Bn	H	Me	NH2	2-Nap	C		479 (M <sup>+</sup> +1)
N-h-248	NA	N-h-247		2CF3Bn	H	H	NH2	2-Nap	C		465 (M <sup>+</sup> +1)
N-h-249	ND1	N-h-67		2CF3Bn	H	Me	NH2	5-Ind	C		468 (M <sup>+</sup> +1)
N-h-250	NA	N-h-249		2CF3Bn	H	H	NH2	5-Ind	C		454 (M <sup>+</sup> +1)
N-h-251	ND1	N-h-69		2CF3Bn	H	Me	NH2	1Me-5-Ind	C		482 (M <sup>+</sup> +1)
N-h-252	NA	N-h-251		2CF3Bn	H	H	NH2	1Me-5-Ind	C		468 (M <sup>+</sup> +1)
N-h-253	ND1	N-h-71		2CF3Bn	H	Me	NH2	5-1HIdz	C		469 (M <sup>+</sup> +1)
N-h-254	NA	N-h-253		2CF3Bn	H	H	NH2	5-1HIdz	C		455 (M <sup>+</sup> +1)
N-h-255	ND1	N-h-73		2-TF	H	Me	NH2	5-Ind	C		406 (M <sup>+</sup> +1)
N-h-256	NA	N-h-255		2-TF	H	H	NH2	5-Ind	C		392 (M <sup>+</sup> +1)
N-h-257	ND1	N-h-75		2-TF	H	Me	NH2	1Me-5-Ind	C		420 (M <sup>+</sup> +1)
N-h-258	NA	N-h-257		2-TF	H	H	NH2	1Me-5-Ind	C		406 (M <sup>+</sup> +1)
N-h-259	ND1	N-h-77		2-TF	H	Me	NH2	1Me-5-1HIdz	C		421 (M <sup>+</sup> +1)
N-h-260	NA	N-h-259		2-TF	H	H	NH2	1Me-5-1HIdz	C		407 (M <sup>+</sup> +1)
N-h-261	ND1	N-h-79		3-TF	H	Me	NH2	5-Ind	C		406 (M <sup>+</sup> +1)
N-h-262	NA	N-h-261		3-TF	H	H	NH2	5-Ind	C		392 (M <sup>+</sup> +1)
N-h-263	ND1	N-h-81		3-TF	H	Me	NH2	1Me-5-Ind	C		420 (M <sup>+</sup> +1)
N-h-264	NA	N-h-263		3-TF	H	H	NH2	1Me-5-Ind	C		406 (M <sup>+</sup> +1)
N-h-265	ND1	N-h-83		3-TF	H	Me	NH2	5-1HIdz	C		407 (M <sup>+</sup> +1)
N-h-266	NA	N-h-265		3-TF	H	H	NH2	5-1HIdz	C		393 (M <sup>+</sup> +1)
N-h-267	ND1	N-h-85		3-TF	H	Me	NH2	1Me-5-1HIdz	C		421 (M <sup>+</sup> +1)
N-h-268	NA	N-h-267		3-TF	H	H	NH2	1Me-5-1HIdz	C		407 (M <sup>+</sup> +1)
N-h-269	ND1	N-h-87		2-FR	H	Me	NH2	2Nap	C		401 (M <sup>+</sup> +1)
N-h-270	NA	N-h-269		2-FR	H	H	NH2	2Nap	C		387 (M <sup>+</sup> +1)
N-h-271	ND1	N-h-89		2-FR	H	Me	NH2	5-Ind	C		390 (M <sup>+</sup> +1)
N-h-272	NA	N-h-271		2-FR	H	H	NH2	5-Ind	C		376 (M <sup>+</sup> +1)
N-h-273	ND1	N-h-91		2-FR	H	Me	NH2	1Me-5-1HIdz	C		405 (M <sup>+</sup> +1)
N-h-274	NA	N-h-273		2-FR	H	H	NH2	1Me-5-1HIdz	C		391 (M <sup>+</sup> +1)



Table-N-H-7

Exp.	Syn	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									method	RTime	Mass
N-h-275	ND1	N-h-93		Bn	Me	Me	NH2	2-Nap	C		425 (M <sup>+</sup> +1)
N-h-276	NA	N-h-275		Bn	Me	H	NH2	2-Nap	C		411 (M <sup>+</sup> +1)
N-h-277	ND1	N-h-95		Bn	Me	Me	NH2	5-Ind	C		414 (M <sup>+</sup> +1)
N-h-278	NA	N-h-277		Bn	Me	H	NH2	5-Ind	C		400 (M <sup>+</sup> +1)
N-h-279	ND1	N-h-97		Bn	Me	Me	NH2	1Me-5-Ind	C		428 (M <sup>+</sup> +1)
N-h-280	NA	N-h-279		Bn	Me	H	NH2	1Me-5-Ind	C		414 (M <sup>+</sup> +1)
N-h-281	ND1	N-h-99		Bn	Me	Me	NH2	5-1HIdz	C		415 (M <sup>+</sup> +1)
N-h-282	NA	N-h-281		Bn	Me	H	NH2	5-1HIdz	C		401 (M <sup>+</sup> +1)
N-h-283	ND1	N-h-101		Bn	Me	Me	NH2	1Me-5-1HIdz	C		429 (M <sup>+</sup> +1)
N-h-284	NA	N-h-283		Bn	Me	H	NH2	1Me-5-1HIdz	C		415 (M <sup>+</sup> +1)
N-h-285	ND1	N-h-103		Bn	Me	Me	NH2	3-Qu	C		426 (M <sup>+</sup> +1)
N-h-286	NA	N-h-285		Bn	Me	H	NH2	3-Qu	C		412 (M <sup>+</sup> +1)
N-h-287	ND1	N-h-105		Bn	Me	Me	NH2	6-Qu	C		426 (M <sup>+</sup> +1)
N-h-288	NA	N-h-287		Bn	Me	H	NH2	6-Qu	C		412 (M <sup>+</sup> +1)
N-h-289	ND1	N-h-107		Bn	Me	Me	NH2	6-IQ	C		426 (M <sup>+</sup> +1)
N-h-290	NA	N-h-289		Bn	Me	H	NH2	6-IQ	C		412 (M <sup>+</sup> +1)
N-h-291	ND1	N-h-109		4FBn	Me	Me	NH2	2-Nap	C		443 (M <sup>+</sup> +1)
N-h-292	NA	N-h-291		4FBn	Me	H	NH2	2-Nap	C		429 (M <sup>+</sup> +1)
N-h-293	ND1	N-h-111		4FBn	Me	Me	NH2	5-Ind	C		432 (M <sup>+</sup> +1)
N-h-294	NA	N-h-293		4FBn	Me	H	NH2	5-Ind	C		418 (M <sup>+</sup> +1)
N-h-295	ND1	N-h-113		4FBn	Me	Me	NH2	1Me-5-Ind	C		446 (M <sup>+</sup> +1)
N-h-296	NA	N-h-295		4FBn	Me	H	NH2	1Me-5-Ind	C		432 (M <sup>+</sup> +1)
N-h-297	ND1	N-h-115		4FBn	Me	Me	NH2	5-1HIdz	C		433 (M <sup>+</sup> +1)
N-h-298	NA	N-h-297		4FBn	Me	H	NH2	5-1HIdz	C		419 (M <sup>+</sup> +1)
N-h-299	ND1	N-h-117		4FBn	Me	Me	NH2	1Me-5-1HIdz	C		447 (M <sup>+</sup> +1)
N-h-300	NA	N-h-299		4FBn	Me	H	NH2	1Me-5-1HIdz	C		433 (M <sup>+</sup> +1)
N-h-301	ND1	N-h-119		2FBn	Me	Me	NH2	2-Nap	C		443 (M <sup>+</sup> +1)
N-h-302	NA	N-h-301		2FBn	Me	H	NH2	2-Nap	C		429 (M <sup>+</sup> +1)
N-h-303	ND1	N-h-121		2FBn	Me	Me	NH2	1Me-5-Ind	C		446 (M <sup>+</sup> +1)
N-h-304	NA	N-h-303		2FBn	Me	H	NH2	1Me-5-Ind	C		432 (M <sup>+</sup> +1)
N-h-305	ND1	N-h-123		2FBn	Me	Me	NH2	5-1HIdz	C		433 (M <sup>+</sup> +1)
N-h-306	NA	N-h-305		2FBn	Me	H	NH2	5-1HIdz	C		419 (M <sup>+</sup> +1)
N-h-307	ND1	N-h-125		2FBn	Me	Me	NH2	1Me-5-1HIdz	C		447 (M <sup>+</sup> +1)
N-h-308	NA	N-h-307		2FBn	Me	H	NH2	1Me-5-1HIdz	C		433 (M <sup>+</sup> +1)
N-h-309	ND1	N-h-127		3FBn	Me	Me	NH2	2-Nap	C		443 (M <sup>+</sup> +1)
N-h-310	NA	N-h-309		3FBn	Me	H	NH2	2-Nap	C		429 (M <sup>+</sup> +1)
N-h-311	ND1	N-h-129		3FBn	Me	Me	NH2	1Me-5-Ind	C		446 (M <sup>+</sup> +1)
N-h-312	NA	N-h-311		3FBn	Me	H	NH2	1Me-5-Ind	C		432 (M <sup>+</sup> +1)
N-h-313	ND1	N-h-131		3FBn	Me	Me	NH2	5-1HIdz	C		433 (M <sup>+</sup> +1)
N-h-314	NA	N-h-313		3FBn	Me	H	NH2	5-1HIdz	C		419 (M <sup>+</sup> +1)
N-h-315	ND1	N-h-133		3FBn	Me	Me	NH2	1Me-5-1HIdz	C		447 (M <sup>+</sup> +1)
N-h-316	NA	N-h-315		3FBn	Me	H	NH2	1Me-5-1HIdz	C		433 (M <sup>+</sup> +1)
N-h-317	ND1	N-h-135		2,3DFBn	Me	Me	NH2	2-Nap	C		461 (M <sup>+</sup> +1)
N-h-318	NA	N-h-317		2,3DFBn	Me	H	NH2	2-Nap	C		447 (M <sup>+</sup> +1)
N-h-319	ND1	N-h-137		2,3DFBn	Me	Me	NH2	1Me-5-Ind	C		464 (M <sup>+</sup> +1)
N-h-320	NA	N-h-319		2,3DFBn	Me	H	NH2	1Me-5-Ind	C		450 (M <sup>+</sup> +1)

Table-N-H-8

Exp.	Syn	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									method	RTime	Mass
N-h-321	ND1	N-h-139		2,3DFBn	Me	Me	NH2	5-1HIdz	C		451 (M <sup>+</sup> +1)
N-h-322	NA	N-h-321		2,3DFBn	Me	H	NH2	5-1HIdz	C		437 (M <sup>+</sup> +1)
N-h-323	ND1	N-h-141		2,3DFBn	Me	Me	NH2	1Me-5-1HIdz	C		465 (M <sup>+</sup> +1)
N-h-324	NA	N-h-323		2,3DFBn	Me	H	NH2	1Me-5-1HIdz	C		451 (M <sup>+</sup> +1)
N-h-325	ND1	N-h-143		3,4DFBn	Me	Me	NH2	2-Nap	C		461 (M <sup>+</sup> +1)
N-h-326	NA	N-h-325		3,4DFBn	Me	H	NH2	2-Nap	C		447 (M <sup>+</sup> +1)
N-h-327	ND1	N-h-145		3,4DFBn	Me	Me	NH2	1Me-5-Ind	C		464 (M <sup>+</sup> +1)
N-h-328	NA	N-h-327		3,4DFBn	Me	H	NH2	1Me-5-Ind	C		450 (M <sup>+</sup> +1)
N-h-329	ND1	N-h-147		3,4DFBn	Me	Me	NH2	1Me-5-1HIdz	C		465 (M <sup>+</sup> +1)
N-h-330	NA	N-h-329		3,4DFBn	Me	H	NH2	1Me-5-1HIdz	C		451 (M <sup>+</sup> +1)
N-h-331	ND1	N-h-149		4PhBn	Me	Me	NH2	2-Nap	C		501 (M <sup>+</sup> +1)
N-h-332	NA	N-h-331		4PhBn	Me	H	NH2	2-Nap	C		487 (M <sup>+</sup> +1)
N-h-333	ND1	N-h-151		4PhBn	Me	Me	NH2	5-Ind	C		490 (M <sup>+</sup> +1)
N-h-334	NA	N-h-333		4PhBn	Me	H	NH2	5-Ind	C		476 (M <sup>+</sup> +1)
N-h-335	ND1	N-h-153		4PhBn	Me	Me	NH2	1Me-5-Ind	C		504 (M <sup>+</sup> +1)
N-h-336	NA	N-h-335		4PhBn	Me	H	NH2	1Me-5-Ind	C		490 (M <sup>+</sup> +1)
N-h-337	ND1	N-h-155		4PhBn	Me	Me	NH2	1Me-5-1HIdz	C		505 (M <sup>+</sup> +1)
N-h-338	NA	N-h-337		4PhBn	Me	H	NH2	1Me-5-1HIdz	C		491 (M <sup>+</sup> +1)
N-h-339	ND1	N-h-157		2CF3Bn	Me	Me	NH2	2-Nap	C		493 (M <sup>+</sup> +1)
N-h-340	NA	N-h-339		2CF3Bn	Me	H	NH2	2-Nap	C		479 (M <sup>+</sup> +1)
N-h-341	ND1	N-h-159		2CF3Bn	Me	Me	NH2	5-Ind	C		482 (M <sup>+</sup> +1)
N-h-342	NA	N-h-341		2CF3Bn	Me	H	NH2	5-Ind	C		468 (M <sup>+</sup> +1)
N-h-343	ND1	N-h-161		2CF3Bn	Me	Me	NH2	1Me-5-Ind	C		496 (M <sup>+</sup> +1)
N-h-344	NA	N-h-343		2CF3Bn	Me	H	NH2	1Me-5-Ind	C		482 (M <sup>+</sup> +1)
N-h-345	ND1	N-h-163		2CF3Bn	Me	Me	NH2	5-1HIdz	C		483 (M <sup>+</sup> +1)
N-h-346	NA	N-h-345		2CF3Bn	Me	H	NH2	5-1HIdz	C		469 (M <sup>+</sup> +1)
N-h-347	ND1	N-h-165		2-TF	Me	Me	NH2	2-Nap	C		431 (M <sup>+</sup> +1)
N-h-348	NA	N-h-347		2-TF	Me	H	NH2	2-Nap	C		417 (M <sup>+</sup> +1)
N-h-349	ND1	N-h-167		2-TF	Me	Me	NH2	1Me-5-Ind	C		434 (M <sup>+</sup> +1)
N-h-350	NA	N-h-349		2-TF	Me	H	NH2	1Me-5-Ind	C		420 (M <sup>+</sup> +1)
N-h-351	ND1	N-h-169		2-TF	Me	Me	NH2	1Me-5-1HIdz	C		435 (M <sup>+</sup> +1)
N-h-352	NA	N-h-351		2-TF	Me	H	NH2	1Me-5-1HIdz	C		421 (M <sup>+</sup> +1)
N-h-353	ND1	N-h-171		3-TF	Me	Me	NH2	2-Nap	C		431 (M <sup>+</sup> +1)
N-h-354	NA	N-h-353		3-TF	Me	H	NH2	2-Nap	C		417 (M <sup>+</sup> +1)
N-h-355	ND1	N-h-173		3-TF	Me	Me	NH2	5-Ind	C		420 (M <sup>+</sup> +1)
N-h-356	NA	N-h-355		3-TF	Me	H	NH2	5-Ind	C		406 (M <sup>+</sup> +1)
N-h-357	ND1	N-h-175		3-TF	Me	Me	NH2	1Me-5-Ind	C		434 (M <sup>+</sup> +1)
N-h-358	NA	N-h-357		3-TF	Me	H	NH2	1Me-5-Ind	C		420 (M <sup>+</sup> +1)
N-h-359	ND1	N-h-177		3-TF	Me	Me	NH2	1Me-5-1HIdz	C		435 (M <sup>+</sup> +1)
N-h-360	NA	N-h-359		3-TF	Me	H	NH2	1Me-5-1HIdz	C		421 (M <sup>+</sup> +1)
N-h-361	ND1	N-h-179		2-FR	Me	Me	NH2	2-Nap	C		415 (M <sup>+</sup> +1)
N-h-362	NA	N-h-361		2-FR	Me	H	NH2	2-Nap	C		401 (M <sup>+</sup> +1)
N-h-363	ND1	N-h-181		2-FR	Me	Me	NH2	5-Ind	C		404 (M <sup>+</sup> +1)
N-h-364	NA	N-h-363		2-FR	Me	H	NH2	5-Ind	C		390 (M <sup>+</sup> +1)
N-h-365	ND1	N-h-183		2-FR	Me	Me	NH2	1Me-5-1HIdz	C		419 (M <sup>+</sup> +1)
N-h-366	NA	N-h-365		2-FR	Me	H	NH2	1Me-5-1HIdz	C		405 (M <sup>+</sup> +1)

Table-N-H-9

Exp.	Syn	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									method	RTime	Mass
N-h-367	NN1	N-h-275	CH <sub>3</sub> I	Bn	Me	Me	NHMe	2-Nap	C		439 (M <sup>+</sup> +1)
N-h-368	NA	N-h-367		Bn	Me	H	NHMe	2-Nap	C		425 (M <sup>+</sup> +1)
N-h-369	NN1	N-h-279	CH <sub>3</sub> I	Bn	Me	Me	NHMe	1Me-5-Ind	C		442 (M <sup>+</sup> +1)
N-h-370	NA	N-h-369		Bn	Me	H	NHMe	1Me-5-Ind	C		428 (M <sup>+</sup> +1)
N-h-371	NN1	N-h-283	CH <sub>3</sub> I	Bn	Me	Me	NHMe	1Me-5-1HIdz	C		443 (M <sup>+</sup> +1)
N-h-372	NA	N-h-371		Bn	Me	H	NHMe	1Me-5-1HIdz	C		429 (M <sup>+</sup> +1)
N-h-373	NN1	N-h-285	CH <sub>3</sub> I	Bn	Me	Me	NHMe	3-Qu	C		440 (M <sup>+</sup> +1)
N-h-374	NA	N-h-373		Bn	Me	H	NHMe	3-Qu	C		426 (M <sup>+</sup> +1)
N-h-375	NN1	N-h-289	CH <sub>3</sub> I	Bn	Me	Me	NHMe	6-IQ	C		440 (M <sup>+</sup> +1)
N-h-376	NA	N-h-375		Bn	Me	H	NHMe	6-IQ	C		426 (M <sup>+</sup> +1)
N-h-377	NN1	N-h-291	CH <sub>3</sub> I	4FBn	Me	Me	NHMe	2-Nap	C		457 (M <sup>+</sup> +1)
N-h-378	NA	N-h-377		4FBn	Me	H	NHMe	2-Nap	C		443 (M <sup>+</sup> +1)
N-h-379	NN1	N-h-295	CH <sub>3</sub> I	4FBn	Me	Me	NHMe	1Me-5-Ind	C		460 (M <sup>+</sup> +1)
N-h-380	NA	N-h-379		4FBn	Me	H	NHMe	1Me-5-Ind	C		446 (M <sup>+</sup> +1)
N-h-381	NN1	N-h-299	CH <sub>3</sub> I	4FBn	Me	Me	NHMe	1Me-5-1HIdz	C		461 (M <sup>+</sup> +1)
N-h-382	NA	N-h-381		4FBn	Me	H	NHMe	1Me-5-1HIdz	C		447 (M <sup>+</sup> +1)
N-h-383	NN1	N-h-301	CH <sub>3</sub> I	2FBn	Me	Me	NHMe	2-Nap	C		457 (M <sup>+</sup> +1)
N-h-384	NA	N-h-383		2FBn	Me	H	NHMe	2-Nap	C		443 (M <sup>+</sup> +1)
N-h-385	NN1	N-h-303	CH <sub>3</sub> I	2FBn	Me	Me	NHMe	1Me-5-Ind	C		460 (M <sup>+</sup> +1)
N-h-386	NA	N-h-385		2FBn	Me	H	NHMe	1Me-5-Ind	C		446 (M <sup>+</sup> +1)
N-h-387	NN1	N-h-307	CH <sub>3</sub> I	2FBn	Me	Me	NHMe	1Me-5-1HIdz	C		461 (M <sup>+</sup> +1)
N-h-388	NA	N-h-387		2FBn	Me	H	NHMe	1Me-5-1HIdz	C		447 (M <sup>+</sup> +1)
N-h-389	NN1	N-h-309	CH <sub>3</sub> I	3FBn	Me	Me	NHMe	2-Nap	C		457 (M <sup>+</sup> +1)
N-h-390	NA	N-h-389		3FBn	Me	H	NHMe	2-Nap	C		443 (M <sup>+</sup> +1)
N-h-391	NN1	N-h-311	CH <sub>3</sub> I	3FBn	Me	Me	NHMe	1Me-5-Ind	C		460 (M <sup>+</sup> +1)
N-h-392	NA	N-h-391		3FBn	Me	H	NHMe	1Me-5-Ind	C		446 (M <sup>+</sup> +1)
N-h-393	NN1	N-h-317	CH <sub>3</sub> I	2,3DFBn	Me	Me	NHMe	2-Nap	C		475 (M <sup>+</sup> +1)
N-h-394	NA	N-h-393		2,3DFBn	Me	H	NHMe	2-Nap	C		461 (M <sup>+</sup> +1)
N-h-395	NN1	N-h-323	CH <sub>3</sub> I	2,3DFBn	Me	Me	NHMe	1Me-5-1HIdz	C		479 (M <sup>+</sup> +1)
N-h-396	NA	N-h-395		2,3DFBn	Me	H	NHMe	1Me-5-1HIdz	C		465 (M <sup>+</sup> +1)
N-h-397	NN1	N-h-327	CH <sub>3</sub> I	3,4DFBn	Me	Me	NHMe	1Me-5-Ind	C		478 (M <sup>+</sup> +1)
N-h-398	NA	N-h-397		3,4DFBn	Me	H	NHMe	1Me-5-Ind	C		464 (M <sup>+</sup> +1)
N-h-399	NN1	N-h-331	CH <sub>3</sub> I	4PhBn	Me	Me	NHMe	2-Nap	C		515 (M <sup>+</sup> +1)
N-h-400	NA	N-h-399		4PhBn	Me	H	NHMe	2-Nap	C		501 (M <sup>+</sup> +1)
N-h-401	NN1	N-h-337	CH <sub>3</sub> I	4PhBn	Me	Me	NHMe	1Me-5-1HIdz	C		519 (M <sup>+</sup> +1)
N-h-402	NA	N-h-401		4PhBn	Me	H	NHMe	1Me-5-1HIdz	C		505 (M <sup>+</sup> +1)
N-h-403	NN1	N-h-339	CH <sub>3</sub> I	2CF3Bn	Me	Me	NHMe	2-Nap	C		507 (M <sup>+</sup> +1)
N-h-404	NA	N-h-403		2CF3Bn	Me	H	NHMe	2-Nap	C		493 (M <sup>+</sup> +1)
N-h-405	NN1	N-h-343	CH <sub>3</sub> I	2CF3Bn	Me	Me	NHMe	1Me-5-Ind	C		510 (M <sup>+</sup> +1)
N-h-406	NA	N-h-405		2CF3Bn	Me	H	NHMe	1Me-5-Ind	C		496 (M <sup>+</sup> +1)
N-h-407	NN1	N-h-347	CH <sub>3</sub> I	2-TF	Me	Me	NHMe	2-Nap	C		445 (M <sup>+</sup> +1)
N-h-408	NA	N-h-407		2-TF	Me	H	NHMe	2-Nap	C		431 (M <sup>+</sup> +1)
N-h-409	NN1	N-h-357	CH <sub>3</sub> I	3-TF	Me	Me	NHMe	1Me-5-Ind	C		448 (M <sup>+</sup> +1)
N-h-410	NA	N-h-409		3-TF	Me	H	NHMe	1Me-5-Ind	C		434 (M <sup>+</sup> +1)
N-h-411	NN1	N-h-365	CH <sub>3</sub> I	2-FR	Me	Me	NHMe	1Me-5-1HIdz	C		433 (M <sup>+</sup> +1)
N-h-412	NA	N-h-411		2-FR	Me	H	NHMe	1Me-5-1HIdz	C		419 (M <sup>+</sup> +1)

Table-N-H-10

Exp.	Syn	SM1	SM2	Rz	Ry	Y	Zx	AR	LCMS		
									method	RTime	Mass
N-h-413	NN2	N-h-275	CH <sub>3</sub> I	Bn	Me	Me	NMe2	2-Nap	C		453 (M <sup>+</sup> +1)
N-h-414	NA	N-h-413		Bn	Me	H	NMe2	2-Nap	C		439 (M <sup>+</sup> +1)
N-h-415	NN2	N-h-279	CH <sub>3</sub> I	Bn	Me	Me	NMe2	1Me-5-Ind	C		456 (M <sup>+</sup> +1)
N-h-416	NA	N-h-415		Bn	Me	H	NMe2	1Me-5-Ind	C		442 (M <sup>+</sup> +1)
N-h-417	NN2	N-h-283	CH <sub>3</sub> I	Bn	Me	Me	NMe2	1Me-5-1HIdz	C		457 (M <sup>+</sup> +1)
N-h-418	NA	N-h-417		Bn	Me	H	NMe2	1Me-5-1HIdz	C		443 (M <sup>+</sup> +1)
N-h-419	NN2	N-h-285	CH <sub>3</sub> I	Bn	Me	Me	NMe2	3-Qu	C		454 (M <sup>+</sup> +1)
N-h-420	NA	N-h-419		Bn	Me	H	NMe2	3-Qu	C		440 (M <sup>+</sup> +1)
N-h-421	NN2	N-h-289	CH <sub>3</sub> I	Bn	Me	Me	NMe2	6-IQ	C		454 (M <sup>+</sup> +1)
N-h-422	NA	N-h-421		Bn	Me	H	NMe2	6-IQ	C		440 (M <sup>+</sup> +1)
N-h-423	NN2	N-h-291	CH <sub>3</sub> I	4FBn	Me	Me	NMe2	2-Nap	C		471 (M <sup>+</sup> +1)
N-h-424	NA	N-h-423		4FBn	Me	H	NMe2	2-Nap	C		457 (M <sup>+</sup> +1)
N-h-425	NN2	N-h-295	CH <sub>3</sub> I	4FBn	Me	Me	NMe2	1Me-5-Ind	C		474 (M <sup>+</sup> +1)
N-h-426	NA	N-h-425		4FBn	Me	H	NMe2	1Me-5-Ind	C		460 (M <sup>+</sup> +1)
N-h-427	NN2	N-h-299	CH <sub>3</sub> I	4FBn	Me	Me	NMe2	1Me-5-1HIdz	C		475 (M <sup>+</sup> +1)
N-h-428	NA	N-h-427		4FBn	Me	H	NMe2	1Me-5-1HIdz	C		461 (M <sup>+</sup> +1)
N-h-429	NN2	N-h-301	CH <sub>3</sub> I	2FBn	Me	Me	NMe2	2-Nap	C		471 (M <sup>+</sup> +1)
N-h-430	NA	N-h-429		2FBn	Me	H	NMe2	2-Nap	C		457 (M <sup>+</sup> +1)
N-h-431	NN2	N-h-303	CH <sub>3</sub> I	2FBn	Me	Me	NMe2	1Me-5-Ind	C		474 (M <sup>+</sup> +1)
N-h-432	NA	N-h-431		2FBn	Me	H	NMe2	1Me-5-Ind	C		460 (M <sup>+</sup> +1)
N-h-433	NN2	N-h-307	CH <sub>3</sub> I	2FBn	Me	Me	NMe2	1Me-5-1HIdz	C		475 (M <sup>+</sup> +1)
N-h-434	NA	N-h-433		2FBn	Me	H	NMe2	1Me-5-1HIdz	C		461 (M <sup>+</sup> +1)
N-h-435	NN2	N-h-309	CH <sub>3</sub> I	3FBn	Me	Me	NMe2	2-Nap	C		471 (M <sup>+</sup> +1)
N-h-436	NA	N-h-435		3FBn	Me	H	NMe2	2-Nap	C		457 (M <sup>+</sup> +1)
N-h-437	NN2	N-h-311	CH <sub>3</sub> I	3FBn	Me	Me	NMe2	1Me-5-Ind	C		474 (M <sup>+</sup> +1)
N-h-438	NA	N-h-437		3FBn	Me	H	NMe2	1Me-5-Ind	C		460 (M <sup>+</sup> +1)
N-h-439	NN2	N-h-317	CH <sub>3</sub> I	2,3DFBn	Me	Me	NMe2	2-Nap	C		489 (M <sup>+</sup> +1)
N-h-440	NA	N-h-439		2,3DFBn	Me	H	NMe2	2-Nap	C		475 (M <sup>+</sup> +1)
N-h-441	NN2	N-h-323	CH <sub>3</sub> I	2,3DFBn	Me	Me	NMe2	1Me-5-1HIdz	C		493 (M <sup>+</sup> +1)
N-h-442	NA	N-h-441		2,3DFBn	Me	H	NMe2	1Me-5-1HIdz	C		479 (M <sup>+</sup> +1)
N-h-443	NN2	N-h-327	CH <sub>3</sub> I	3,4DFBn	Me	Me	NMe2	1Me-5-Ind	C		492 (M <sup>+</sup> +1)
N-h-444	NA	N-h-443		3,4DFBn	Me	H	NMe2	1Me-5-Ind	C		478 (M <sup>+</sup> +1)
N-h-445	NN2	N-h-331	CH <sub>3</sub> I	4PhBn	Me	Me	NMe2	2-Nap	C		529 (M <sup>+</sup> +1)
N-h-446	NA	N-h-445		4PhBn	Me	H	NMe2	2-Nap	C		515 (M <sup>+</sup> +1)
N-h-447	NN2	N-h-337	CH <sub>3</sub> I	4PhBn	Me	Me	NMe2	1Me-5-1HIdz	C		533 (M <sup>+</sup> +1)
N-h-448	NA	N-h-447		4PhBn	Me	H	NMe2	1Me-5-1HIdz	C		519 (M <sup>+</sup> +1)
N-h-449	NN2	N-h-339	CH <sub>3</sub> I	2CF3Bn	Me	Me	NMe2	2-Nap	C		521 (M <sup>+</sup> +1)
N-h-450	NA	N-h-449		2CF3Bn	Me	H	NMe2	2-Nap	C		507 (M <sup>+</sup> +1)
N-h-451	NN2	N-h-343	CH <sub>3</sub> I	2CF3Bn	Me	Me	NMe2	1Me-5-Ind	C		524 (M <sup>+</sup> +1)
N-h-452	NA	N-h-451		2CF3Bn	Me	H	NMe2	1Me-5-Ind	C		510 (M <sup>+</sup> +1)
N-h-453	NN2	N-h-347	CH <sub>3</sub> I	2-TF	Me	Me	NMe2	2-Nap	C		459 (M <sup>+</sup> +1)
N-h-454	NA	N-h-453		2-TF	Me	H	NMe2	2-Nap	C		445 (M <sup>+</sup> +1)
N-h-455	NN2	N-h-357	CH <sub>3</sub> I	3-TF	Me	Me	NMe2	1Me-5-Ind	C		462 (M <sup>+</sup> +1)
N-h-456	NA	N-h-455		3-TF	Me	H	NMe2	1Me-5-Ind	C		448 (M <sup>+</sup> +1)
N-h-457	NN2	N-h-365	CH <sub>3</sub> I	2-FR	Me	Me	NMe2	1Me-5-1HIdz	C		447 (M <sup>+</sup> +1)
N-h-458	NA	N-h-457		2-FR	Me	H	NMe2	1Me-5-1HIdz	C		433 (M <sup>+</sup> +1)

[Examples N-i-1 to N-i-138]

Typical examples of the compounds of the present invention that can be obtained by reacting and treating corresponding starting compounds using any of the methods described in the present specification are shown in Table-N-I-1 to Table-N-I-8. In the tables, the compound numbers are mentioned in the columns indicated as "Exp.". In the tables, used methods among the aforementioned

synthesis methods are shown in the columns of "Syn" with symbols, the starting compounds 1 are mentioned in the columns of "SM1", and the starting compounds 2 are mentioned in the columns of "SM2".

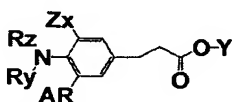


Table-N-I-1

Exp.	Syn	SM1	SM2	NRzRy	Y	Zx	AR	LCMS		
								method	RTime	Mass
N-i-1	NB1	Int.n-111	BRA1		Me	NO2	2-Nap	C		405 (M <sup>+</sup> +1)
N-i-2	NA	N-i-1			H	NO2	2-Nap	C		391 (M <sup>+</sup> +1)
N-i-3	NB1	Int.n-111	BRA2		Me	NO2	5-1Ind	C		394 (M <sup>+</sup> +1)
N-i-4	NA	N-i-3			H	NO2	5-1Ind	C		380 (M <sup>+</sup> +1)
N-i-5	NB1	Int.n-111	BRA3		Me	NO2	1Me-5-Ind	C		408 (M <sup>+</sup> +1)
N-i-6	NA	N-i-5			H	NO2	1Me-5-Ind	C		394 (M <sup>+</sup> +1)
N-i-7	NB1	Int.n-111	BRA5		Me	NO2	5-1Hldz	C		395 (M <sup>+</sup> +1)
N-i-8	NA	N-i-7			H	NO2	5-1Hldz	C		381 (M <sup>+</sup> +1)
N-i-9	NB1	Int.n-111	BRA6		Me	NO2	1Me-5-1Hldz	C		409 (M <sup>+</sup> +1)
N-i-10	NA	N-i-9			H	NO2	1Me-5-1Hldz	C		395 (M <sup>+</sup> +1)
N-i-11	NB1	Int.n-111	BRA9		Me	NO2	5-Bzt	C		412 (M <sup>+</sup> +1)
N-i-12	NA	N-i-11			H	NO2	5-Bzt	C		398 (M <sup>+</sup> +1)
N-i-13	NB1	Int.n-111	BRA 10		Me	NO2	3-Qu	C		406 (M <sup>+</sup> +1)
N-i-14	NA	N-i-13			H	NO2	3-Qu	C		392 (M <sup>+</sup> +1)
N-i-15	NB1	Int.n-111	BRA 11		Me	NO2	6-Qu	C		406 (M <sup>+</sup> +1)
N-i-16	NA	N-i-15			H	NO2	6-Qu	C		392 (M <sup>+</sup> +1)
N-i-17	NB1	Int.n-112	BRA1		Me	NO2	2-Nap	C		421 (M <sup>+</sup> +1)
N-i-18	NA	N-i-17			H	NO2	2-Nap	C		407 (M <sup>+</sup> +1)
N-i-19	NB1	Int.n-112	BRA2		Me	NO2	5-1Ind	C		410 (M <sup>+</sup> +1)
N-i-20	NA	N-i-19			H	NO2	5-1Ind	C		396 (M <sup>+</sup> +1)
N-i-21	NB1	Int.n-112	BRA3		Me	NO2	1Me-5-Ind	C		424 (M <sup>+</sup> +1)
N-i-22	NA	N-i-21			H	NO2	1Me-5-Ind	C		410 (M <sup>+</sup> +1)

Table-N-I-2























Exp.	Syn	SM1	SM2	NRzRy	Y	Zx	AR	LCMS		
								method	RTime	Mass
N-i-23	NB1	Int.n-112	BRA5		Me	NO2	5-1HIdz	C		411 (M <sup>+</sup> +1)
N-i-24	NA	N-i-23			H	NO2	5-1HIdz	C		397 (M <sup>+</sup> +1)
N-i-25	NB1	Int.n-112	BRA6		Me	NO2	1Me-5-1HIdz	C		425 (M <sup>+</sup> +1)
N-i-26	NA	N-i-25			H	NO2	1Me-5-1HIdz	C		411 (M <sup>+</sup> +1)
N-i-27	NB1	Int.n-113	BRA1		Me	NO2	2-Nap	C		419 (M <sup>+</sup> +1)
N-i-28	NA	N-i-27			H	NO2	2-Nap	C		405 (M <sup>+</sup> +1)
N-i-29	NB1	Int.n-113	BRA2		Me	NO2	5-1Ind	C		408 (M <sup>+</sup> +1)
N-i-30	NA	N-i-29			H	NO2	5-1Ind	C		394 (M <sup>+</sup> +1)
N-i-31	NB1	Int.n-113	BRA3		Me	NO2	1Me-5-Ind	C		422 (M <sup>+</sup> +1)
N-i-32	NA	N-i-31			H	NO2	1Me-5-Ind	C		408 (M <sup>+</sup> +1)
N-i-33	NB1	Int.n-113	BRA5		Me	NO2	5-1HIdz	C		409 (M <sup>+</sup> +1)
N-i-34	NA	N-i-33			H	NO2	5-1HIdz	C		395 (M <sup>+</sup> +1)
N-i-35	NB1	Int.n-113	BRA6		Me	NO2	1Me-5-1HIdz	C		423 (M <sup>+</sup> +1)
N-i-36	NA	N-i-35			H	NO2	1Me-5-1HIdz	C		409 (M <sup>+</sup> +1)
N-i-37	NB1	Int.n-113	BRA11		Me	NO2	6-Qu	C		420 (M <sup>+</sup> +1)
N-i-38	NA	N-i-37			H	NO2	6-Qu	C		406 (M <sup>+</sup> +1)
N-i-39	NB1	Int.n-114	BRA1		Me	NO2	2-Nap	C		433 (M <sup>+</sup> +1)
N-i-40	NA	N-i-39			H	NO2	2-Nap	C		419 (M <sup>+</sup> +1)
N-i-41	NB1	Int.n-114	BRA3		Me	NO2	1Me-5-Ind	C		437 (M <sup>+</sup> +1)
N-i-42	NA	N-i-41			H	NO2	1Me-5-Ind	C		423 (M <sup>+</sup> +1)
N-i-43	NB1	Int.n-114	BRA5		Me	NO2	5-1HIdz	C		423 (M <sup>+</sup> +1)
N-i-44	NA	N-i-43			H	NO2	5-1HIdz	C		409 (M <sup>+</sup> +1)

Table-N-I-3

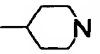





















Exp.	Syn	SM1	SM2	NRzRy	Y	Zx	AR	LCMS		
								method	RTime	Mass
N-i-45	NB1	Int.n-114	BRA6		Me	NO2	1Me-5-1HIdz	C		437 (M <sup>+</sup> +1)
N-i-46	NA	N-i-45			H	NO2	1Me-5-1HIdz	C		423 (M <sup>+</sup> +1)
N-i-47	NB1	Int.n-115	BRA3		Me	NO2	1Me-5-Ind	C		436 (M <sup>+</sup> +1)
N-i-48	NA	N-i-47			H	NO2	1Me-5-Ind	C		422 (M <sup>+</sup> +1)
N-i-49	NB1	Int.n-115	BRA5		Me	NO2	5-1HIdz	C		423 (M <sup>+</sup> +1)
N-i-50	NA	N-i-49			H	NO2	5-1HIdz	C		409 (M <sup>+</sup> +1)
N-i-51	NB1	Int.n-115	BRA6		Me	NO2	1Me-5-1HIdz	C		437 (M <sup>+</sup> +1)
N-i-52	NA	N-i-51			H	NO2	1Me-5-1HIdz	C		423 (M <sup>+</sup> +1)
N-i-53	ND1	N-i-1			Me	NH2	2-Nap	C		375 (M <sup>+</sup> +1)
N-i-54	NA	N-i-53			H	NH2	2-Nap	C		361 (M <sup>+</sup> +1)
N-i-55	ND1	N-i-3			Me	NH2	5-1Ind	C		364 (M <sup>+</sup> +1)
N-i-56	NA	N-i-55			H	NH2	5-1Ind	C		350 (M <sup>+</sup> +1)
N-i-57	ND1	N-i-5			Me	NH2	1Me-5-Ind	C		378 (M <sup>+</sup> +1)
N-i-58	NA	N-i-57			H	NH2	1Me-5-Ind	C		364 (M <sup>+</sup> +1)
N-i-59	ND1	N-i-7			Me	NH2	5-1HIdz	C		365 (M <sup>+</sup> +1)
N-i-60	NA	N-i-59			H	NH2	5-1HIdz	C		351 (M <sup>+</sup> +1)
N-i-61	ND1	N-i-9			Me	NH2	1Me-5-1HIdz	C		379 (M <sup>+</sup> +1)
N-i-62	NA	N-i-61			H	NH2	1Me-5-1HIdz	C		365 (M <sup>+</sup> +1)
N-i-63	ND1	N-i-11			Me	NH2	5-Bzt	C		382 (M <sup>+</sup> +1)
N-i-64	NA	N-i-63			H	NH2	5-Bzt	C		368 (M <sup>+</sup> +1)
N-i-65	ND1	N-i-13			Me	NH2	3-Qu	C		376 (M <sup>+</sup> +1)
N-i-66	NA	N-i-65			H	NH2	3-Qu	C		362 (M <sup>+</sup> +1)

Table-N-I-4























Exp.	Syn	SM1	SM2	NRzRy	Y	Zx	AR	LCMS		
								method	RTime	Mass
N-i-67	ND1	N-i-15			Me	NH2	6-Qu	C		376 (M <sup>+</sup> +1)
N-i-68	NA	N-i-67			H	NH2	6-Qu	C		362 (M <sup>+</sup> +1)
N-i-69	ND1	N-i-17			Me	NH2	2-Nap	C		391 (M <sup>+</sup> +1)
N-i-70	NA	N-i-69			H	NH2	2-Nap	C		377 (M <sup>+</sup> +1)
N-i-71	ND1	N-i-19			Me	NH2	5-1Ind	C		380 (M <sup>+</sup> +1)
N-i-72	NA	N-i-71			H	NH2	5-1Ind	C		366 (M <sup>+</sup> +1)
N-i-73	ND1	N-i-21			Me	NH2	1Me-5-Ind	C		394 (M <sup>+</sup> +1)
N-i-74	NA	N-i-73			H	NH2	1Me-5-Ind	C		380 (M <sup>+</sup> +1)
N-i-75	ND1	N-i-23			Me	NH2	5-1HIdz	C		381 (M <sup>+</sup> +1)
N-i-76	NA	N-i-75			H	NH2	5-1HIdz	C		367 (M <sup>+</sup> +1)
N-i-77	ND1	N-i-25			Me	NH2	1Me-5-1HIdz	C		395 (M <sup>+</sup> +1)
N-i-78	NA	N-i-77			H	NH2	1Me-5-1HIdz	C		381 (M <sup>+</sup> +1)
N-i-79	ND1	N-i-27			Me	NH2	2-Nap	C		389 (M <sup>+</sup> +1)
N-i-80	NA	N-i-79			H	NH2	2-Nap	C		375 (M <sup>+</sup> +1)
N-i-81	ND1	N-i-29			Me	NH2	5-1Ind	C		378 (M <sup>+</sup> +1)
N-i-82	NA	N-i-81			H	NH2	5-1Ind	C		364 (M <sup>+</sup> +1)
N-i-83	ND1	N-i-31			Me	NH2	1Me-5-Ind	C		392 (M <sup>+</sup> +1)
N-i-84	NA	N-i-83			H	NH2	1Me-5-Ind	C		378 (M <sup>+</sup> +1)
N-i-85	ND1	N-i-33			Me	NH2	5-1HIdz	C		379 (M <sup>+</sup> +1)
N-i-86	NA	N-i-85			H	NH2	5-1HIdz	C		365 (M <sup>+</sup> +1)
N-i-87	ND1	N-i-35			Me	NH2	1Me-5-1HIdz	C		393 (M <sup>+</sup> +1)
N-i-88	NA	N-i-87			H	NH2	1Me-5-1HIdz	C		379 (M <sup>+</sup> +1)



Table-N-I-5























Exp.	Syn	SM1	SM2	NRzRy	Y	Zx	AR	LCMS		
								method	RTime	Mass
N-i-89	ND1	N-i-37			Me	NH2	6-Qu	C		390 (M <sup>+</sup> +1)
N-i-90	NA	N-i-89			H	NH2	6-Qu	C		376 (M <sup>+</sup> +1)
N-i-91	ND1	N-i-39			Me	NH2	2-Nap	C		403 (M <sup>+</sup> +1)
N-i-92	NA	N-i-91			H	NH2	2-Nap	C		389 (M <sup>+</sup> +1)
N-i-93	ND1	N-i-41			Me	NH2	1Me-5-Ind	C		406 (M <sup>+</sup> +1)
N-i-94	NA	N-i-93			H	NH2	1Me-5-Ind	C		392 (M <sup>+</sup> +1)
N-i-95	ND1	N-i-43			Me	NH2	5-1Idz	C		393 (M <sup>+</sup> +1)
N-i-96	NA	N-i-95			H	NH2	5-1Idz	C		379 (M <sup>+</sup> +1)
N-i-97	ND1	N-i-45			Me	NH2	1Me-5-1HIdz	C		407 (M <sup>+</sup> +1)
N-i-98	NA	N-i-97			H	NH2	1Me-5-1HIdz	C		393 (M <sup>+</sup> +1)
N-i-99	ND1	N-i-47			Me	NH2	1Me-5-Ind	C		406 (M <sup>+</sup> +1)
N-i-100	NA	N-i-99			H	NH2	1Me-5-Ind	C		392 (M <sup>+</sup> +1)
N-i-101	ND1	N-i-49			Me	NH2	5-1Idz	C		393 (M <sup>+</sup> +1)
N-i-102	NA	N-i-101			H	NH2	5-1Idz	C		379 (M <sup>+</sup> +1)
N-i-103	ND1	N-i-51			Me	NH2	1Me-5-1HIdz	C		407 (M <sup>+</sup> +1)
N-i-104	NA	N-i-103			H	NH2	1Me-5-1HIdz	C		393 (M <sup>+</sup> +1)
N-i-105	NN1	N-i-53	CH <sub>3</sub> I		Me	NHMe	2-Nap	C		389 (M <sup>+</sup> +1)
N-i-106	NA	N-i-105			H	NHMe	2-Nap	C		375 (M <sup>+</sup> +1)
N-i-107	NN1	N-i-57	CH <sub>3</sub> I		Me	NHMe	1Me-5-Ind	C		392 (M <sup>+</sup> +1)
N-i-108	NA	N-i-107			H	NHMe	1Me-5-Ind	C		378 (M <sup>+</sup> +1)
N-i-109	NN1	N-i-61	CH <sub>3</sub> I		Me	NHMe	1Me-5-1HIdz	C		393 (M <sup>+</sup> +1)
N-i-110	NA	N-i-109			H	NHMe	1Me-5-1HIdz	C		379 (M <sup>+</sup> +1)

Table-N-I-6



















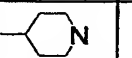
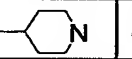


Exp.	Syn	SM1	SM2	NRzRy	Y	Zx	AR	LCMS		
								method	RTime	Mass
N-i-111	NN1	N-i-63	CH <sub>3</sub> I		Me	NHMe	5-Bzt	C		396 (M <sup>+</sup> +1)
N-i-112	NA	N-i-111			H	NHMe	5-Bzt	C		382 (M <sup>+</sup> +1)
N-i-113	NN1	N-i-65	CH <sub>3</sub> I		Me	NHMe	3-Qu	C		390 (M <sup>+</sup> +1)
N-i-114	NA	N-i-113			H	NHMe	3-Qu	C		376 (M <sup>+</sup> +1)
N-i-115	NN1	N-i-67	CH <sub>3</sub> I		Me	NHMe	6-Qu	C		390 (M <sup>+</sup> +1)
N-i-116	NA	N-i-115			H	NHMe	6-Qu	C		376 (M <sup>+</sup> +1)
N-i-117	NN1	N-i-69	CH <sub>3</sub> I		Me	NHMe	2-Nap	C		405 (M <sup>+</sup> +1)
N-i-118	NA	N-i-117			H	NHMe	2-Nap	C		391 (M <sup>+</sup> +1)
N-i-119	NN1	N-i-73	CH <sub>3</sub> I		Me	NHMe	1Me-5-Ind	C		408 (M <sup>+</sup> +1)
N-i-120	NA	N-i-119			H	NHMe	1Me-5-Ind	C		394 (M <sup>+</sup> +1)
N-i-121	NN1	N-i-77	CH <sub>3</sub> I		Me	NHMe	1Me-5-1HIdz	C		409 (M <sup>+</sup> +1)
N-i-122	NA	N-i-121			H	NHMe	1Me-5-1HIdz	C		395 (M <sup>+</sup> +1)
N-i-123	NN1	N-i-79	CH <sub>3</sub> I		Me	NHMe	2-Nap	C		403 (M <sup>+</sup> +1)
N-i-124	NA	N-i-123			H	NHMe	2-Nap	C		389 (M <sup>+</sup> +1)
N-i-125	NN1	N-i-83	CH <sub>3</sub> I		Me	NHMe	1Me-5-Ind	C		406 (M <sup>+</sup> +1)
N-i-126	NA	N-i-125			H	NHMe	1Me-5-Ind	C		392 (M <sup>+</sup> +1)
N-i-127	NN1	N-i-87	CH <sub>3</sub> I		Me	NHMe	1Me-5-1HIdz	C		407 (M <sup>+</sup> +1)
N-i-128	NA	N-i-127			H	NHMe	1Me-5-1HIdz	C		393 (M <sup>+</sup> +1)
N-i-129	NN1	N-i-91	CH <sub>3</sub> I		Me	NHMe	2-Nap	C		417 (M <sup>+</sup> +1)
N-i-130	NA	N-i-129			H	NHMe	2-Nap	C		403 (M <sup>+</sup> +1)
N-i-131	NN1	N-i-93	CH <sub>3</sub> I		Me	NHMe	1Me-5-Ind	C		420 (M <sup>+</sup> +1)
N-i-132	NA	N-i-131			H	NHMe	1Me-5-Ind	C		406 (M <sup>+</sup> +1)

Table-N-I-7































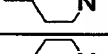

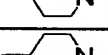
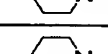

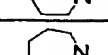




Exp.	Syn	SM1	SM2	NRzRy	Y	Zx	AR	LCMS		
								method	RTime	Mass
N-i-133	NN1	N-i-97	CH <sub>3</sub> I		Me	NHMe	1Me-5-1HIdz	C		421 (M <sup>+</sup> +1)
N-i-134	NA	N-i-133			H	NHMe	1Me-5-1HIdz	C		407 (M <sup>+</sup> +1)
N-i-135	NN1	N-i-99	CH <sub>3</sub> I		Me	NHMe	1Me-5-Ind	C		420 (M <sup>+</sup> +1)
N-i-136	NA	N-i-135			H	NHMe	1Me-5-Ind	C		406 (M <sup>+</sup> +1)
N-i-137	NN1	N-i-103	CH <sub>3</sub> I		Me	NHMe	1Me-5-1HIdz	C		421 (M <sup>+</sup> +1)
N-i-138	NA	N-i-137			H	NHMe	1Me-5-1HIdz	C		407 (M <sup>+</sup> +1)
N-i-139	NN2	N-i-53	CH <sub>3</sub> I		Me	NMe <sub>2</sub>	2-Nap	C		403 (M <sup>+</sup> +1)
N-i-140	NA	N-i-139			H	NMe <sub>2</sub>	2-Nap	C		389 (M <sup>+</sup> +1)
N-i-141	NN2	N-i-57	CH <sub>3</sub> I		Me	NMe <sub>2</sub>	1Me-5-Ind	C		406 (M <sup>+</sup> +1)
N-i-142	NA	N-i-141			H	NMe <sub>2</sub>	1Me-5-Ind	C		392 (M <sup>+</sup> +1)
N-i-143	NN2	N-i-61	CH <sub>3</sub> I		Me	NMe <sub>2</sub>	1Me-5-1HIdz	C		407 (M <sup>+</sup> +1)
N-i-144	NA	N-i-143			H	NMe <sub>2</sub>	1Me-5-1HIdz	C		393 (M <sup>+</sup> +1)
N-i-145	NN2	N-i-63	CH <sub>3</sub> I		Me	NMe <sub>2</sub>	5-Bzt	C		410 (M <sup>+</sup> +1)
N-i-146	NA	N-i-145			H	NMe <sub>2</sub>	5-Bzt	C		396 (M <sup>+</sup> +1)
N-i-147	NN2	N-i-65	CH <sub>3</sub> I		Me	NMe <sub>2</sub>	3-Qu	C		404 (M <sup>+</sup> +1)
N-i-148	NA	N-i-147			H	NMe <sub>2</sub>	3-Qu	C		390 (M <sup>+</sup> +1)
N-i-149	NN2	N-i-67	CH <sub>3</sub> I		Me	NMe <sub>2</sub>	6-Qu	C		404 (M <sup>+</sup> +1)
N-i-150	NA	N-i-149			H	NMe <sub>2</sub>	6-Qu	C		390 (M <sup>+</sup> +1)
N-i-151	NN2	N-i-69	CH <sub>3</sub> I		Me	NMe <sub>2</sub>	2-Nap	C		419 (M <sup>+</sup> +1)
N-i-152	NA	N-i-151			H	NMe <sub>2</sub>	2-Nap	C		405 (M <sup>+</sup> +1)
N-i-153	NN2	N-i-73	CH <sub>3</sub> I		Me	NMe <sub>2</sub>	1Me-5-Ind	C		422 (M <sup>+</sup> +1)
N-i-154	NA	N-i-153			H	NMe <sub>2</sub>	1Me-5-Ind	C		408 (M <sup>+</sup> +1)

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Exp.	Syn	SM1	SM2	NRzRy	Y	Zx	AR	LCMS		
								method	RTime	Mass
N-i-121	NN2	N-i-77	CH <sub>3</sub> I		Me	NMe2	1Me-5-1HIdz	C		423 (M <sup>+</sup> +1)
N-i-122	NA	N-i-121			H	NMe2	1Me-5-1HIdz	C		409 (M <sup>+</sup> +1)
N-i-123	NN2	N-i-79	CH <sub>3</sub> I		Me	NMe2	2-Nap	C		417 (M <sup>+</sup> +1)
N-i-124	NA	N-i-123			H	NMe2	2-Nap	C		403 (M <sup>+</sup> +1)
N-i-125	NN2	N-i-83	CH <sub>3</sub> I		Me	NMe2	1Me-5-Ind	C		420 (M <sup>+</sup> +1)
N-i-126	NA	N-i-125			H	NMe2	1Me-5-Ind	C		406 (M <sup>+</sup> +1)
N-i-127	NN2	N-i-87	CH <sub>3</sub> I		Me	NMe2	1Me-5-1HIdz	C		421 (M <sup>+</sup> +1)
N-i-128	NA	N-i-127			H	NMe2	1Me-5-1HIdz	C		407 (M <sup>+</sup> +1)
N-i-129	NN2	N-i-91	CH <sub>3</sub> I		Me	NMe2	2-Nap	C		431 (M <sup>+</sup> +1)
N-i-130	NA	N-i-129			H	NMe2	2-Nap	C		417 (M <sup>+</sup> +1)
N-i-131	NN2	N-i-93	CH <sub>3</sub> I		Me	NMe2	1Me-5-Ind	C		434 (M <sup>+</sup> +1)
N-i-132	NA	N-i-131			H	NMe2	1Me-5-Ind	C		420 (M <sup>+</sup> +1)
N-i-133	NN2	N-i-97	CH <sub>3</sub> I		Me	NMe2	1Me-5-1HIdz	C		435 (M <sup>+</sup> +1)
N-i-134	NA	N-i-133			H	NMe2	1Me-5-1HIdz	C		421 (M <sup>+</sup> +1)
N-i-135	NN2	N-i-99	CH <sub>3</sub> I		Me	NMe2	2-Nap	C		431 (M <sup>+</sup> +1)
N-i-136	NA	N-i-135			H	NMe2	2-Nap	C		417 (M <sup>+</sup> +1)
N-i-137	NN2	N-i-103	CH <sub>3</sub> I		Me	NMe2	1Me-5-1HIdz	C		435 (M <sup>+</sup> +1)
N-i-138	NA	N-i-137			H	NMe2	1Me-5-1HIdz	C		421 (M <sup>+</sup> +1)

## [Test Examples]

1. Suppressing Action on PGE<sub>2</sub> production from IL-1  $\beta$  -stimulated MG-63 cells

## (1) Method for measurement

An action of suppressing PGE<sub>2</sub> production caused by interleukin (IL) 1  $\beta$  as an inflammatory stimulant was studied by the following method. Cells of MG-63, which is a human osteosarcoma cell line (purchased from Dainippon Pharmaceutical), were suspended in EMEM medium (GIBCO) containing 10% fetal

bovine serum (BioFluid), and then inoculated to each well of 96-well culture plate at a density of  $2 \times 10^4$  cells/well and cultured overnight. The medium was changed to EMEM medium containing 0.5% fetal bovine serum, and then a test compound was added to each well. Human interleukin- $1\beta$  (ENDOGEN) was further added as an inflammatory stimulant at a final concentration of 1 ng/ml. The cells were further cultured for 18 hours. Then, the culture supernatant was collected, and the PGE<sub>2</sub> concentration in the culture supernatant was measured by using EIA kit (CAYMAN). By using a well which was not added with the stimulant as a negative control and a well which was added only with the stimulant as a positive control, suppression ratio on PGE<sub>2</sub> production was calculated from the produced amount of PGE<sub>2</sub> in the well added with the test compound using the following equation.

[Equation 1]

$$\text{PGE}_2 \text{ production suppression ratio} = [1 - (C - B)/(A - B)] \times 100$$

A: PGE<sub>2</sub> production amount of positive control

B: PGE<sub>2</sub> production amount of negative control

C: PGE<sub>2</sub> production amount in well added with test compound

Further, cytotoxicity of the compounds was studied by using the cells after the collection of the supernatant according to the methylene blue uptake method. Specifically, the cells remained after the collection of the supernatant were fixed with glutaraldehyde and stained with a 0.05% methylene blue solution, then methylene blue taken up by the cells was extracted with 0.3 N hydrochloric acid, and absorbance of the extract was measured at 670 nm. The absorbance of the well of the aforementioned positive control was taken as 100%, and a test compound that gave absorbance in well of less than 80% was judged to be positive in cytotoxicity.

## (2) Measurement results

The test compounds (Compound Nos. G-1 to G-121, H-1 to H-32, J-1 to J-92,

K-1 to K-40, L-1 to L-95, M-1 to M-32, N-1 to N-74, P-1 to P-50, Q-1 to Q-52, S-1 to S-73, T-1 to T-61, U-1 to U-18, V-1 to V-109, and W-1 to W-13) suppressed the PGE<sub>2</sub> production caused by IL-1 $\beta$  by 50% or more at 1.0  $\mu$  M. Moreover, all the test compounds did not exhibit cytotoxicity at that concentration.

The test compounds (Compound Nos. Ca-1 to Ca-203) suppressed the PGE<sub>2</sub> production caused by IL-1 $\beta$  by 50% or more at 1.0  $\mu$  M. None of the test compounds exhibited cytotoxicity at that concentration.

The test compounds (Compound Nos. S-a-1 to S-a-24, S-b-1 to S-b-138, and S-c-1 to S-c-138) suppressed the PGE<sub>2</sub> production caused by IL-1 $\beta$  by 50% or more at 1.0  $\mu$  M. None of the test compounds exhibited cytotoxicity at that concentration.

Further, the test compounds (Compound Nos. N-a-1 to N-a-142, N-b-1 to N-b-182, N-c-1 to N-c-64, N-d-1 to N-d-74, N-e-1 to N-e-186 and N-g-1 to N-g-44) suppressed the PGE<sub>2</sub> production caused by IL-1 $\beta$  by 50% or more at 1.0  $\mu$  M. None of the test compounds exhibited cytotoxicity at that concentration.

Therefore, the novel substituted phenylalkanoic acid derivatives or salts thereof according to the present invention are useful as agents for suppressing inflammatory prostaglandin production.

## 2. Suppressing action on PGD<sub>2</sub> and LTB<sub>4</sub> production from IgE-stimulated RBL-2H3 cells

### (1) Method for measurement

Suppressing action on PGD<sub>2</sub> and LTB<sub>4</sub> production caused by IgE as an allergic stimulant was investigated by the following method. Cells of RBL-2H3, which is a rat mastocytoma cell line (purchased from ATCC), were suspended in DEMEM medium (GIBCO) containing 10% fetal bovine serum (BioFluid), inoculated to each well of 48-well culture plate at a density of  $2 \times 10^4$  cells/well and cultured overnight. Then, IgE antiserum directed to dinitrophenylated BSA

(hereinafter abbreviated as "DNP-BSA") was further added to each well, and the cells were cultured for 30 minutes. Then, the medium was changed to DEMEM medium containing 0.5% fetal bovine serum, a test compound was added to each well, and DNP-BSA was further added at a final concentration of 100 ng/ml as a stimulant. Ten minutes after the stimulant was added, the culture supernatant was collected, and the PGD<sub>2</sub> concentration and LTB<sub>4</sub> concentration in the culture supernatant were measured by using EIA kit (CAYMAN). By using a well which was not added with the stimulant as a negative control and a well which was added only with the stimulant as a positive control, suppressing ratios on mediator production were calculated from the production amounts of the mediators in the well added with the test compound using the following equation 2.

[Equation 2]

$$\text{PGD}_2 \text{ or LTB}_4 \text{ production suppression ratio} = [1 - (C - B)/(A - B)] \times 100$$

A: PGD<sub>2</sub> or LTB<sub>4</sub> production amount of positive control

B: PGD<sub>2</sub> or LTB<sub>4</sub> production amount of negative control

C: PGD<sub>2</sub> or LTB<sub>4</sub> production amount in well added with test compound

Cytotoxicity of the compounds was studied in the same manner as those described above, by using the cells after the collection of the supernatant according to the methylene blue uptake method.

## (2) Measurement results

Representative compounds of the objective Compounds (I) described in the specification suppressed the PGD<sub>2</sub> and LTB<sub>4</sub> production caused by IgE stimulation by 50% or more at 1.0  $\mu$  M. Moreover, all the test compounds did not exhibit cytotoxicity at that concentration. Thus, the novel substituted phenylalkanoic acid derivatives or salts thereof according to the present invention exhibit suppressing action on the allergic prostaglandin and leukotriene production, and are useful as suppressing agents for the production thereof.

### 3. Suppressing effect on mouse zymosan-stimulated footpad edema reaction

#### (1) Method for measurement

A suppressing effect on footpad edema caused by zymosan as an inflammatory stimulant was studied by the following method. Groups of ICR female mice (6- to 7-week old) each consisting of eight mice were used for the test. A test compound was suspended or dissolved in purified water containing 0.5% methylcellulose and orally administered to the test animals at 0.1 to 500 mg/10 ml/kg. To the control group, purified water containing 0.5% methylcellulose was administered in a similar manner, which was not added with a test compound. One hour after the administration of the test compound, 0.02 ml of a suspension of zymosan suspended in physiological saline (Otsuka Pharmaceutical) at 1 mg/ml was subcutaneously administered to right hind leg footpad of each mouse. One and two hours after the administration of the zymosan suspension, volume of the right hind leg footpad was measured by using an apparatus for measuring a volume of mouse hind leg footpad edema (Unicom). A difference of the volume of footpad measured above and the footpad volume before the administration of the test compound measured beforehand was regarded as a volume of the edema.

For the volume of the edema at 1 hour or 2 hours after the zymosan administration, a graph was prepared by indicating time in abscissa and the edema volume in ordinate, and an edema volume AUC (area under the curve) was obtained up to 2 hours by calculation using the following equation.

[Equation 3]

$$\text{Edema volume AUC } (\mu\text{l}\cdot\text{hour}) = 1/2 \times 1 \times A + 1 \times (A + B)/2$$

A: Edema volume 1 hour after zymosan administration

B: Edema volume 2 hour after zymosan administration

A suppression ratio on edema of test compound was obtained by calculation using the following equation.



[Equation 4]

Edema suppression ratio (%) =  $[1 - B/A] \times 100$

A: Edema volume AUC of positive control

B: Edema volume AUC of test compound administered group

## (2) Measurement results

Representative compounds of the objective Compounds (I) described in the specification more effectively suppressed footpad edema caused by subcutaneous administration of zymosan compared with the positive control group by oral administration at 0.1 to 500 mg/kg.

Therefore, the novel substituted phenylalkanoic acid derivatives or salts thereof according to the present invention exhibit a suppressing action on footpad edema caused by zymosan as an inflammatory stimulant, and thus they are useful as agents for prophylactic and/or therapeutic drugs for inflammatory diseases.

## 4. Suppressing effect on mouse IgE-stimulated footpad edema reaction

### (1) Method for measurement

Suppression on footpad edema caused by IgE antibody as an allergic stimulant was studied by the following method. Groups of C57BL/6 male mice (9- to 11-week old) each consisting of five mice were used for the test. Anti-DNP-BSA IgE serum was subcutaneously administered in a volume of 20  $\mu$ l to right hind leg footpad of each mouse one day before the test. A test compound was suspended or dissolved in purified water containing 0.5% methylcellulose and orally administered to the test animals at 0.1 to 500 mg/10 ml/kg. To the control group, purified water containing 0.5% methylcellulose was administered in a similar manner, which was not added with any test compound. Two hours after the administration of the test compound, 0.2 ml of a solution of DNP-BSA dissolved in physiological saline (Otsuka Pharmaceutical) at 2.5  $\mu$ g/ml was intravenously administered. The thickness of right hind leg footpad was measured by using a

digital thickness gauge (MITSUTOYO) 10, 15, 20, and 30 minutes after the administration of DNP-BSA. A difference of the thickness of footpad measured above and the thickness before the administration of the test compound measured beforehand was regarded as a thickness of edema.

For the thickness of the edema at 10, 15, 20 and 30 minutes after the DNP-BSA administration, a graph was prepared indicating time in abscissa and the edema thickness in ordinate, and edema thickness AUC up to 2 hours was obtained by calculation according to the following equation.

[Equation 5]

$$\begin{aligned} \text{Edema thickness AUC (mm} \cdot \text{minute)} = & 1/2 \times 10 \times A + 5 \times (A + B)/2 \\ & + 5 \times (B + C)/2 + 10 \times (C + D)/2 \end{aligned}$$

A: Edema thickness 10 minutes after DNP-BSA administration

B: Edema thickness 15 minutes after DNP-BSA administration

C: Edema thickness 20 minutes after DNP-BSA administration

D: Edema thickness 30 minutes after DNP-BSA administration

A suppressing ratio on edema of a test compound was obtained by calculation in accordance with the following equation.

[Equation 6]

$$\text{Edema suppression ratio (\%)} = [1 - B/A] \times 100.$$

A: Edema thickness AUC of positive control

B: Edema thickness AUC of test compound administered group

## (2) Measurement results

Representative compounds of the objective Compounds (I) described in the specification suppressed the footpad edema caused by IgE stimulation, i.e., footpad edema observed when DNP-BSA was administered to the mice sensitized with the anti-DNP-BSA IgE serum, compared with the positive control group by oral administration of 0.1 to 500 mg/kg.

Therefore, the novel substituted phenylalkanoic acid derivatives or salts thereof according to the present invention exhibit suppressing action on footpad edema caused by IgE antibody, which is an allergic stimulant, and thus they are useful as prophylactic and/or therapeutic drugs for allergic diseases.

#### 5. Suppressing effect on mouse acetic acid writhing reaction

##### (1) Method for measurement

A suppressing effect on acetic acid writhing reaction, which is an acute pain model, was studied by the following method. Groups of ICR female mice (6-week old) each consisting of eight mice were used for the test. A test compound was suspended or dissolved in purified water containing 0.5% methylcellulose and orally administered to the test animals at 0.1 to 500 mg/10 ml/kg. To the control group, purified water containing 0.5% methylcellulose was administered in a similar manner, which was not added with any test compound. One hour after the administration of the test compound, 0.9% aqueous acetic acid was intraperitoneally administered to the mice in a volume of 5 ml/kg, and number of writhing reactions during 15 minutes immediately after the administration of acetic acid was counted. Suppression ratio relative to the control group was obtained by calculation according to the following equation.

[Equation 7]

Writhing suppression ratio (%) =  $[1 - B/A] \times 100$

A: Writhing number of positive control group

B: Writhing number of test compound administered group

##### (2) Measurement results

The representative compounds of the objective Compounds (I) described in the specification suppressed writhing caused by administration of aqueous acetic acid compared with the positive control group at oral administration of 0.1 to 500 mg/kg.

It has been elucidated that a writhing reaction caused by intraperitoneal administration of acetic acid is caused due to production of prostaglandin [Matsumoto et al., European Journal of Pharmacology (Eur. J. Pharmacol), 1998, vol. 352, p.47; Ueno et al., Biochemical Pharmacology (Biochem. Pharmacol), 2001, vol. 15, p.157].

Therefore, the novel substituted phenylalkanoic acid derivatives or salts thereof according to the present invention are useful as prophylactic and/or therapeutic agents for acute pain caused by prostaglandins.

#### 6. Prophylactic and therapeutic effects for rat adjuvant arthritis

##### (1) Method for measurement

A suppressing effect on footpad edema observed in rat adjuvant arthritis, which is a disease model of rheumatoid arthritis as being one of autoimmune diseases and also a chronic inflammatory disease, was studied by the following method. Groups of Lewis female rats (8-week old) each consisting of six mice were used for the test. The test animals were immunized by subcutaneously administering, to right hind leg footpads, 50  $\mu$ l of liquid paraffin containing 10 mg/ml of M. tuberculosis H37 RA (DIFCO) as an adjuvant. A test compound was suspended or dissolved in purified water containing 0.5% methylcellulose and orally administered to the test animals at 0.1 to 500 mg/5 ml/kg. The test compound was administered twice a day for 14 days, from the 12th day after the immunization. To the control group, purified water containing 0.5% methylcellulose was administered in a similar manner, which was not added with any test compound. Every 2 or 3 days after the administration of adjuvant, volume of left hind leg footpad, which was not administered with the adjuvant, was measured by using an apparatus for measuring a volume of edema of a rat hind leg footpad (Unicom). A suppression ratio on edema was obtained by calculation using the following equation.

[Equation 8]

Edema suppression ratio (%) =  $\{1 - [(D - C)/C]/[(B - A)/A]\} \times 100$

A: Left hind leg footpad volume of positive control immediately before administration of adjuvant

B: Left hind leg footpad volume of positive control on each measurement day

C: Left hind leg footpad volume of test compound administered group immediately before administration of adjuvant

D: Left hind leg footpad volume of test compound administered group on each measurement day

## (2) Measurement results

The representative compounds of the objective Compound (I) described in the specification suppressed footpad edema in adjuvant arthritis compared with the positive control group.

Therefore, the novel substituted phenylalkanoic acid derivatives or salts thereof according to the present invention are useful as agents for prophylactic and/or therapeutic drugs for rheumatoid arthritis and autoimmune diseases.

## 7. Effect on rat pulmonary fibrosis

### (1) Method for measurement

A suppressing effect on pulmonary fibrosis in a bleomycin-induced rat pulmonary fibrosis model, which is a pathological model of pulmonary fibrosis, was studied by the following method. Groups of BN female rats (7-week old) each consisting of seven rats were used for the test. The test animals were anesthetized with ketamine and xylazine, and the tracheae were exposed. Then, a 125  $\mu$ g/0.1 ml solution of bleomycin (Nippon Kayaku) dissolved in physiological saline (Ohtsuka Pharmaceutical Factory) was injected into the tracheae by using a syringe. The negative control group was administered with 0.1 ml of saline into

the tracheae.

Each test compound was suspended or dissolved in purified water containing 0.5% methylcellulose, and orally administered to the test animals at doses of 10, 30, 100 and 300 mg/5 ml/kg. The administration of the test compounds was started from the day of the bleomycin administration and performed once or twice a day for 21 days. The positive control group was administered with purified water containing 0.5% methylcellulose not added with any test compound in a similar manner. On the 21st day after the administration of bleomycin, the rats were sacrificed, and lungs were fixed with neutral buffered formalin to prepare histopathological samples. Staining of the histopathological samples was performed by the Azan method.

The histopathological samples of lungs were examined, and degree of fibrosing was represented with the following scores on the basis of formation of granulation tissues and proliferation of collagen fibers as indicators, i.e., -: no abnormality,  $\pm$ : extremely mild change, +: mild change, ++: moderate change, and +++: significant change.

## (2) Measurement results

The fibrosing score of the negative control group was minus (-), and no pulmonary fibrosing was observed. The median of the fibrosing score of the positive control group was from ++ to +++, and pulmonary fibrosing was observed. The medians of the fibrosing score of the groups of rats administered with the test compounds (Compound Nos. G-2, G-4 and V-40) were from  $\pm$  to +, and thus the fibrosing was milder compared with the positive control group. The median of the fibrosing score of the group administered with the other test compounds (Compound Nos. G118 and V-59) was from  $\pm$  to +, and thus pulmonary fibrosing was milder than that observed in the positive control group. Accordingly, the compounds of the present invention are useful as a prophylactic and/or therapeutic agent for

pulmonary fibrosis, and type 4 PLA<sub>2</sub> inhibitor compounds are useful as a prophylactic and/or therapeutic agent (including a progression-preventing agent) for pulmonary fibrosis.

Further, known cPLA<sub>2</sub> inhibitory compounds, arachidonyl trifluoromethyl ketone, 4-(1-benzhydryl-6-chloro-1H-indol-3-ylmethyl)-3-methoxybenzoic acid, N-{1-[2-(2,4-difluorobenzoyl)benzoyl]-4-tritylsulfanylpyrrolidin-2-ylmethyl}-4-(2,4-dioxothiazolidin-5-ylidenemethyl)benzoic acid amide and 4-{4-[2-(2-[bis(4-chlorophenyl)methoxy]ethylsulfonyl)ethoxy]phenyl}-1,1,1-trifluoro-2-butanone, are intraperitoneally or orally administered in a similar manner. Fibrosing is mild also in the groups administered with these known type 4 PLA<sub>2</sub> inhibitory compounds.

#### Industrial Applicability

The compounds of the present invention have superior suppressing action on prostaglandin production and leukotriene production, and they are useful as active ingredients of medicaments for prophylactic and/or therapeutic treatment of various inflammatory diseases, autoimmune diseases, allergic diseases, pain, fibrosis and the like caused by these lipid mediators.